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(54) Title: ESSENTIAL BACTERIAL GENES AND THEIR USE

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Disclosed are 23 genes, termed "GEP" genes, found in *streptococcus pneumonia*, which are located within operons that are essential for survival. Also disclosed is a related essential gene found in *Bacillus subtilis*. These genes and the polypeptides that they encode, as well as homologs thereof, can be used to identify antibacterial agents for treating bacterial infections such as streptococcal pneumonia.

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ESSENTIAL BACTERIAL GENES AND THEIR USE

Background of the Invention

The invention relates to essential bacterial genes and their use in identifying antibacterial agents.

5 Bacterial infections may be cutaneous, subcutaneous, or systemic. Opportunistic bacterial infections proliferate, especially in patients afflicted with AIDS or other diseases that compromise the immune system. The bacterium *Streptococcus pneumonia* typically infects the respiratory tract and can cause lobar pneumonia, as well as meningitis, sinusitis, and other infections.

Summary of the Invention

10 The invention is based on the discovery of 23 genes in the bacterium *Streptococcus pneumoniae*, and a related gene in the bacterium *Bacillus subtilis*, that are located within operons that are essential for survival. These 23 *Streptococcus* genes are referred to herein as "GEP genes" (which stands for
15 general essential protein); for convenience, the polypeptides encoded by these genes are referred to herein as "GEP polypeptides." Each GEP gene is located within an operon that contains a gene that is essential for survival of *Streptococcus pneumoniae*; the essential gene can be the GEP gene or another gene located within the same operon. Bacterial operons contain several genes that are related, e.g.,
20 with respect to function or biochemical pathway. Transcription of an operon leads to the production of a single transcript in which multiple coding regions are linked. Thus, an operon containing one or more essential genes can be considered an "essential operon," since disruption of expression of one gene located within the operon will interfere with expression of the other genes in the operon. Each coding
25 region of the transcript is separately translated into an individual polypeptide by ribosomes that initiate translation at multiple points along the transcript. Having identified one gene in the operon, one can readily identify and sequence the other genes located within the operon.

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The genes encoding the GEP polypeptides are useful molecular tools for identifying similar genes in pathogenic microorganisms, such as pathogenic strains of *Bacillus*. In addition, the operons containing genes encoding GEP polypeptides, and the polypeptides encoded by such operons, are useful targets for identifying
5 compounds that are inhibitors of the pathogens in which the GEP polypeptides are expressed. Such inhibitors inhibit bacterial growth by being bacteriostatic (e.g., inhibiting reproduction or cell division) or by being bacteriocidal (i.e., by causing cell death).

The invention, therefore, features an isolated polypeptide encoded by a
10 nucleic acid located within an operon encoding a GEP polypeptide, termed gep103, having the amino acid sequence set forth in SEQ ID NO:1, or conservative variations thereof. An isolated operon comprising a nucleic acid encoding gep103 also is included within the invention. In addition, the invention includes an isolated nucleic acid of (a) an operon comprising the sequence of SEQ ID NO:2, as
15 depicted in Fig. 1, or degenerate variants thereof; (b) an operon comprising the sequence of SEQ ID NO:2, or degenerate variants thereof, wherein T is replaced by U; (c) nucleic acids complementary to (a) and (b); and (d) fragments of (a), (b), and (c) that are at least 15 base pairs in length and that hybridize under stringent conditions to genomic DNA encoding the polypeptide of SEQ ID NO:1. As
20 described above for gep103, other nucleic acids and polypeptides encoded by nucleic acids located within operons encoding GEP polypeptides are included within the invention, including: (a) operons comprising the nucleic acids represented by the SEQ ID NOs. listed below, as depicted in the Figures listed below, or degenerate variants thereof; (b) operons comprising the nucleic acids
25 represented by the SEQ ID NOs. listed below, wherein T is replaced by U; (c) nucleic acids complementary to (a) and (b); and (d) fragments of (a), (b), and (c) that are at least 15 base pairs in length and that hybridize under stringent conditions to genomic DNA encoding the polypeptides represented by the SEQ ID NOs. listed below.

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Table 1: GEP nucleic acids and polypeptides

	GEP Nucleic Acid or Polypeptide	Figure No.	SEQ ID No. of Amino Acid Sequence	SEQ ID No. of the Coding Strand of the Nucleic Acid Sequence	SEQ ID No. of the Non-coding Strand of the Nucleic Acid Sequence
5	gep103	1	1	2	3
	gep1119	2	4	5	6
	gep1122	3	7	8	9
	gep1315	4	10	11	12
10	gep1493	5	13	14	15
	gep1507	6	16	17	18
	gep1511	7	19	20	21
	gep1518	8	22	23	24
15	gep1546	9	25	26	27
	gep1551	10	28	29	30
	gep1561	11	31	32	33
	gep1580	12	34	35	36
20	gep1713	13	37	38	39
	gep222	14	40	41	42
	gep2283	15	43	44	45
	gep273	16	46	47	48
25	gep286	17	49	50	51
	gep311	18	52	53	54
	gep3262	19	55	56	57
	gep3387	20	58	59	60
	gep47	21	61	62	63

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GEP Nucleic Acid or Polypeptide	Figure No.	SEQ ID No. of Amino Acid Sequence	SEQ ID No. of the Coding Strand of the Nucleic Acid Sequence	SEQ ID No. of the Non-coding Strand of the Nucleic Acid Sequence
gep61	22	64	65	66
gep76	23	67	68	69

The invention also includes allelic variants (i.e., genes encoding isozymes) of the genes located within operons encoding the GEP polypeptides listed above.

- 5 For example, the invention includes a gene that encodes a GEP polypeptide but which gene includes one or more point mutations, deletions, promotor variants, or splice site variants, provided that the resulting GEP polypeptide functions as a GEP polypeptide (e.g., as determined in a conventional complementation assay).

- Identification of these GEP genes and the determination that they are
- 10 located within operons containing an essential gene allows homologs of the GEP genes to be found in other organisms strains of *Streptococcus*. Also, orthologs of these genes can be identified in other species (e.g., *Bacillus sp.*). While "homologs" are structurally similar genes contained within a species, "orthologs" are functionally equivalent genes from other species (within or outside of a given
- 15 genus, e.g., from *Bacillus subtilis* or *E. coli*). Such homologs and orthologs are expected to be located within operons that are essential for survival. Such homologous and orthologous genes and polypeptides can be used to identify compounds that inhibit the growth of the host organism (e.g., compounds that are bacteriocidal or bacteriostatic against pathogenic strains of the organism).
- 20 Homologous and orthologous genes and polypeptides that are essential for survival can serve as targets for identifying a broad spectrum of antibacterial agents.

An ortholog of gep1493, termed B-yneS, has been identified in *B. subtilis* and is essential for survival of *B. subtilis*. The amino acid sequence (SEQ ID NO: 70), coding sequence (SEQ ID NO:71), and non-coding sequence (SEQ ID NO:72)

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of B-yneS is set forth in Fig. 24. As with the other polypeptides and genes disclosed herein, the B-yneS polypeptide and gene can be used in the methods described herein to identify antibacterial agents.

The term gep103 polypeptide or gene as used herein is intended to include the polypeptide and gene set forth in Fig. 1 herein, as well as homologs of the sequences set forth in Fig. 1. Also encompassed by the term gep103 gene are degenerate variants of the nucleic acid sequence set forth in Fig. 1 (SEQ ID NO:2). Degenerate variants of a nucleic acid sequence exist because of the degeneracy of the amino acid code; thus, those sequences that vary from the sequence represented by SEQ ID NO:2, but which nonetheless encode a gep103 polypeptide are included within the invention. Likewise, because of the similarity in the structures of amino acids, conservative variations (as described herein) can be made in the amino acid sequence of the gep103 polypeptide while retaining the function of the polypeptide (e.g., as determined in a conventional complementation assay). Other gep103 polypeptides and genes identified in additional *Streptococcus* strains may be such conservative variations or degenerate variants of the particular gep103 polypeptide and nucleic acid set forth in Fig. 1 (SEQ ID NOs:1 and 2, respectively). The gep103 polypeptide and gene share at least 80%, e.g., 90%, sequence identity with SEQ ID NOs:1 and 2, respectively. Regardless of the percent sequence identity between the gep103 sequence and the sequence represented by SEQ ID NOs:1 and 2, the gep103 genes and polypeptides encompassed by the invention are able to complement for the lack of gep103 function (e.g., in a temperature-sensitive mutant) in a standard complementation assay. Additional gep103 genes that are identified and cloned from additional *Streptococcus* strains, and pathogenic strains in particular, can be used to produce gep103 polypeptides for use in the various methods described herein, e.g., for identifying antibacterial agents. Likewise, the terms gep1119, gep1122, gep1315, gep1493, gep1507, gep1511, gep1518, gep1546, gep1551, gep1561, gep1580, gep1713, gep222, gep2283, gep273, gep286, gep311, gep3262, gep3387, gep47, gep61, and gep76 encompass homologs, conservative variations, and degenerate variants of the sequences depicted in Figs. 2-23,

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respectively. Such homologs, conservative variations, and degenerate variants also are included within the invention.

Since the various GEP genes described herein have been identified and shown to be located within operons that are essential for survival, the GEP genes
5 and polypeptides encoded by nucleic acid sequences located within operons containing GEP genes and their homologs and orthologs can be used to identify antibacterial agents. More specifically, the polypeptides encoded by nucleic acid sequences located within operons containing GEP genes can be used, separately or together, in assays to identify test compounds that bind to these polypeptides. Such
10 test compounds are expected to be antibacterial agents, in contrast to compounds that do not bind to these GEP polypeptides. As described herein, any of a variety of art-known methods can be used to assay for binding of test compounds to the polypeptides. The invention includes, for example, a method for identifying an antibacterial agent where the method entails: (a) contacting a polypeptide encoded
15 by a nucleic acid sequence located within an operon containing a GEP gene, or homolog or ortholog thereof, with a test compound; (b) detecting binding of the test compound to the polypeptide or homolog or ortholog; and (c) determining whether a test compound that binds to the polypeptide or homolog or ortholog inhibits growth of bacteria, relative to growth of bacteria cultured in the absence of
20 the test compound that binds to the polypeptide or homolog or ortholog, as an indication that the test compound is an antibacterial agent.

In various embodiments, the GEP polypeptide is derived from a non-pathogenic or pathogenic *Streptococcus* strain, such as *Streptococcus pneumoniae*, *Streptococcus pyogenes*, *Streptococcus agalactiae*, *Streptococcus endocarditis*,
25 *Streptococcus faecium*, *Streptococcus sanguis*, *Streptococcus viridans*, and *Streptococcus hemolyticus*. Suitable orthologs of the *Streptococcus* GEP genes can be derived from the bacterium *Bacillus subtilis*. The test compound can be immobilized on a substrate, and binding of the test compound to the polypeptide or homolog or ortholog can be detected as immobilization of the polypeptide or

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homolog or ortholog on the immobilized test compound, e.g., in an immunoassay with an antibody that specifically binds to the polypeptide.

If desired, the test compound can be a test polypeptide (e.g., a polypeptide having a random or predetermined amino acid sequence; or a naturally-occurring or synthetic polypeptide). Alternatively, the test compound can be a nucleic acid, such as a DNA or RNA molecule. In addition, small organic molecules can be tested. The test compound can be a naturally-occurring compound or it can be synthetically produced, if desired. Synthetic libraries, chemical libraries, and the like can be screened to identify compounds that bind to the polypeptides. More generally, binding of test compounds to the polypeptide or homolog or ortholog can be detected either *in vitro* or *in vivo*. Regardless of the source of the test compound, the polypeptides described herein can be used to identify compounds that are bacterioidial or bacteriostatic to a variety of pathogenic or non-pathogenic strains.

In an exemplary method, binding of a test compound to a polypeptide encoded by a nucleic acid located within an operon containing a GEP gene can be detected in a conventional two-hybrid system for detecting protein/protein interactions (e.g., in yeast or mammalian cells). Generally, in such a method, (a) the polypeptide encoded by a nucleic acid located within an operon containing a GEP gene is provided as a fusion protein that includes the polypeptide fused to (i) a transcription activation domain of a transcription factor or (ii) a DNA-binding domain of a transcription factor; (b) the test polypeptide is provided as a fusion protein that includes the test polypeptide fused to (i) a transcription activation domain of a transcription factor or (ii) a DNA-binding domain of a transcription factor; and (c) binding of the test polypeptide to the polypeptide is detected as reconstitution of a transcription factor. Homologs and orthologs of the GEP polypeptides can be used in similar methods. Reconstitution of the transcription factor can be detected, for example, by detecting transcription of a gene that is operably linked to a DNA sequence bound by the DNA-binding domain of the reconstituted transcription factor (See, for example, White, 1996, Proc. Natl. Acad.

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Sci. 93:10001-10003 and references cited therein and Vidal et al., 1996, Proc. Natl. Acad. Sci. 93:10315-10320).

In an alternative method, an isolated operon containing a nucleic acid molecule encoding a GEP polypeptide is used to identify a compound that
5 decreases the expression of a GEP polypeptide *in vivo*. Such compounds can be used as antibacterial agents. To discover such compounds, cells that express a GEP polypeptide are cultured, exposed to a test compound (or a mixture of test compounds), and the level of expression or activity is compared with the level of GEP polypeptide expression or activity in cells that are otherwise identical but that
10 have not been exposed to the test compound(s). Many standard quantitative assays of gene expression can be utilized in this aspect of the invention.

To identify compounds that modulate expression of a GEP polypeptide (or homologous or orthologous sequence), the test compound(s) can be added at varying concentrations to the culture medium of cells that express a GEP
15 polypeptide (or homolog or ortholog), as described herein. Such test compounds can include small molecules (typically, non-protein, non-polysaccharide chemical entities), polypeptides, and nucleic acids. The expression of the GEP polypeptide is then measured, for example, by Northern blot PCR analysis or RNase protection analyses using a nucleic acid molecule of the invention as a probe. The level of
20 expression in the presence of the test molecule, compared with the level of expression in its absence, will indicate whether or not the test molecule alters the expression of the GEP polypeptide. Because the GEP polypeptides are expressed from operons that are essential for survival, test compounds that inhibit the expression and/or function of the GEP polypeptide will inhibit growth of the cells
25 or kill the cells.

Compounds that modulate the expression of the polypeptides of the invention can be identified by carrying out the assays described herein and then measuring the levels of the GEP polypeptides expressed in the cells, e.g., by performing a Western blot analysis using antibodies that bind to a GEP
30 polypeptide.

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- The invention further features methods of identifying from a large group of mutants those strains that have conditional lethal mutations. In general, the gene and corresponding gene product are subsequently identified, although the strains themselves can be used in screening or diagnostic assays. The mechanism(s) of
- 5 action for the identified genes and gene products provide a rational basis for the design of antibacterial therapeutic agents. These antibacterial agents reduce the action of the gene product in a wild type strain, and therefore are useful in treating a subject with that type, or a similarly susceptible type of infection by administering the agent to the subject in a pharmaceutically effective amount.
- 10 Reduction in the action of the gene product includes competitive inhibition of the gene product for the active site of an enzyme or receptor; non-competitive inhibition; disrupting an intracellular cascade path which requires the gene product; binding to the gene product itself, before or after post-translational processing; and acting as a gene product mimetic, thereby down-regulating the activity.
- 15 Therapeutic agents include monoclonal antibodies raised against the gene product.

Furthermore, the presence of the gene sequence in certain cells (e.g., a pathogenic bacterium of the same genus or similar species), and the absence or divergence of the sequence in host cells can be determined, if desired. Therapeutic agents directed toward genes or gene products that are not present in the host have

20 several advantages, including fewer side effects, and lower overall dosage.

- The invention includes pharmaceutical formulations that include a pharmaceutically acceptable excipient and an antibacterial agent identified using the methods described herein. In particular, the invention includes pharmaceutical formulations that contain antibacterial agents that inhibit the growth of, or kill,
- 25 pathogenic *Streptococcus* strains. Such pharmaceutical formulations can be used for treating a *Streptococcus* infection in an organism. Such a method entails administering to the organism a therapeutically effective amount of the pharmaceutical formulation. In particular, such pharmaceutical formulations can be used to treat streptococcal pneumonia in mammals such as humans and
- 30 domesticated mammals (e.g., cows, pigs, dogs, and cats), and in plants. The

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efficacy of such antibacterial agents in humans can be estimated in an animal model system well known to those of skill in the art (e.g., mouse and rabbit model systems).

Also included within the invention are polyclonal and monoclonal antibodies
5 that specifically bind to the various GEP polypeptides described herein (e.g., gep103). Such antibodies can facilitate detection of GEP polypeptides in various *Streptococcus* strains. These antibodies also are useful for detecting binding of a test compound to GEP polypeptides (e.g., using the assays described herein). In addition, monoclonal antibodies that bind to GEP polypeptides are themselves
10 adequate antibacterial agents when administered to a mammal, as such monoclonal antibodies are expected to impede one or more functions of GEP polypeptides.

As used herein, "nucleic acids" encompass both RNA and DNA, including genomic DNA and synthetic (e.g., chemically synthesized) DNA. The nucleic acid can be double-stranded or single-stranded. Where single-stranded, the nucleic acid
15 may be a sense strand or an antisense strand. The nucleic acid may be synthesized using oligonucleotide analogs or derivatives (e.g., inosine or phosphorothioate nucleotides). Such oligonucleotides can be used, for example, to prepare nucleic acids that have altered base-pairing abilities or increased resistance to nucleases.

An "isolated nucleic acid" is a DNA or RNA that is not immediately
20 contiguous with both of the coding sequences with which it is immediately contiguous (one on the 5' end and one on the 3' end) in the naturally occurring genome of the organism from which it is derived. Thus, in one embodiment, an isolated nucleic acid includes some or all of the 5' non-coding (e.g., promoter) sequences that are immediately contiguous to the coding sequence. The term
25 therefore includes, for example, a recombinant DNA that is incorporated into a vector, into an autonomously replicating plasmid or virus, or into the genomic DNA of a prokaryote or eukaryote, or which exists as a separate molecule (e.g., a genomic DNA fragment produced by PCR or restriction endonuclease treatment) independent of other sequences. It also includes a recombinant DNA that is part of
30 a hybrid gene encoding an additional polypeptide sequence. The term "isolated"

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can refer to a nucleic acid or polypeptide that is substantially free of cellular material, viral material, or culture medium (when produced by recombinant DNA techniques), or chemical precursors or other chemicals (when chemically synthesized). Moreover, an "isolated nucleic acid fragment" is a nucleic acid
5 fragment that is not naturally occurring as a fragment and would not be found in the natural state. As used herein, the term "isolated nucleic acid molecule" includes an operon containing a contiguous cluster of linked sequences. "Isolated operons" are those operons that are not naturally occurring and which are not associated with the sequences by which they are normally surrounded in a bacterial genome.

10 A nucleic acid sequence that is "substantially identical" to a GEP nucleotide sequence is at least 80% (e.g., 85%) identical to the nucleotide sequence of the nucleic acid sequences represented by the SEQ ID NOs listed in Table 1, as depicted in Figs. 1-23. For purposes of comparison of nucleic acids, the length of the reference nucleic acid sequence will generally be at least 40 nucleotides, e.g., at
15 least 60 nucleotides or more nucleotides. Sequence identity can be measured using sequence analysis software (e.g., Sequence Analysis Software Package of the Genetics Computer Group, University of Wisconsin Biotechnology Center, 1710 University Avenue, Madison, WI 53705).

The GEP polypeptides useful in practicing the invention include, but are not
20 limited to, recombinant polypeptides and natural polypeptides. Also useful in the invention are nucleic acid sequences that encode forms of GEP polypeptides in which naturally occurring amino acid sequences are altered or deleted. Preferred nucleic acids encode polypeptides that are soluble under normal physiological conditions. Also within the invention are nucleic acids encoding fusion proteins in
25 which a portion of a GEP polypeptide is fused to an unrelated polypeptide (e.g., a marker polypeptide or a fusion partner) to create a fusion protein. For example, the polypeptide can be fused to a hexa-histidine tag to facilitate purification of bacterially expressed polypeptides, or to a hemagglutinin tag to facilitate purification of polypeptides expressed in eukaryotic cells. The invention also
30 includes, for example, isolated polypeptides (and the nucleic acids that encode these

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polypeptides) that include a first portion and a second portion; the first portion includes, e.g., a GEP polypeptide, and the second portion includes an immunoglobulin constant (Fc) region or a detectable marker.

The fusion partner can be, for example, a polypeptide which facilitates
5 secretion, e.g., a secretory sequence. Such a fused polypeptide is typically referred to as a preprotein. The secretory sequence can be cleaved by the host cell to form the mature protein. Also within the invention are nucleic acids that encode a GEP polypeptide fused to a polypeptide sequence to produce an inactive preprotein. Preproteins can be converted into the active form of the protein by removal of the
10 inactivating sequence.

The invention also includes nucleic acids that hybridize, e.g., under stringent hybridization conditions (as defined herein) to all or a portion of the nucleotide sequences represented by the SEQ ID NOs. listed in Table 1, or their complements. The hybridizing portion of the hybridizing nucleic acids is typically at least 15
15 (e.g., 20, 30, or 50) nucleotides in length. The hybridizing portion of the hybridizing nucleic acid is at least 80%, e.g., at least 95%, or at least 98%, identical to the sequence of a portion or all of a nucleic acid encoding a GEP polypeptide or its complement. Hybridizing nucleic acids of the type described herein can be used as a cloning probe, a primer (e.g., a PCR primer), or a
20 diagnostic probe. Nucleic acids that hybridize to the nucleotide sequences represented by the SEQ ID NOs. listed in Table 1 are considered "antisense oligonucleotides." Also included within the invention are ribozymes that inhibit the function of operons containing the GEP genes of the invention, as determined, for example, in a complementation assay.

25 Also useful in the invention are various cells, e.g., transformed host cells, that contain a GEP nucleic acid described herein. A "transformed cell" is a cell into which (or into an ancestor of which) has been introduced, by means of recombinant DNA techniques, a nucleic acid encoding a GEP polypeptide. Both prokaryotic and eukaryotic cells are included, e.g., bacteria, *Streptococcus*, *Bacillus*,
30 and the like.

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Also useful in the invention are genetic constructs (e.g., vectors and plasmids) that include a nucleic acid of the invention which is operably linked to a transcription and/or translation sequence to enable expression, e.g., expression vectors. By "operably linked" is meant that a selected nucleic acid, e.g., a DNA molecule encoding a GEP polypeptide, is positioned adjacent to one or more sequence elements, e.g., a promoter, which directs transcription and/or translation of the sequence such that the sequence elements can control transcription and/or translation of the selected nucleic acid.

The invention also features purified or isolated polypeptides encoded by nucleic acids located within operons containing GEP genes, as listed in Table 1. As used herein, both "protein" and "polypeptide" mean any chain of amino acids, regardless of length or post-translational modification (e.g., glycosylation or phosphorylation). Thus, the terms gep103 polypeptide, gep1119 polypeptide, gep1122 polypeptide, gep1315 polypeptide, gep1493 polypeptide, gep1507 polypeptide, gep1511 polypeptide, gep1518 polypeptide, gep1546 polypeptide, gep1551 polypeptide, gep1561 polypeptide, gep1580 polypeptide, gep1713 polypeptide, gep222 polypeptide, gep2283 polypeptide, gep273 polypeptide, gep286 polypeptide, gep311 polypeptide, gep3262 polypeptide, gep3387 polypeptide, gep47 polypeptide, gep61 polypeptide, and gep76 polypeptide include full-length, naturally occurring gep103, gep1119, gep1122, gep1315, gep1493, gep1507, gep1511, gep1518, gep1546, gep1551, gep1561, gep1580, gep1713, gep222, gep2283, gep273, gep286, gep311, gep3262, gep3387, gep47, gep61, and gep76 proteins, respectively, as well as recombinantly or synthetically produced polypeptides that correspond to the full-length, naturally occurring proteins, or to a portion of the naturally occurring or synthetic polypeptide.

A "purified" or "isolated" compound is a composition that is at least 60% by weight the compound of interest, e.g., a GEP polypeptide or antibody. Preferably the preparation is at least 75% (e.g., at least 90% or 99%) by weight the compound of interest. Purity can be measured by any appropriate standard method, e.g., column chromatography, polyacrylamide gel electrophoresis, or HPLC analysis.

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Preferred GEP polypeptides include a sequence substantially identical to all or a portion of a naturally occurring GEP polypeptide, e.g., including all or a portion of the sequences shown in Figs. 1-23. Polypeptides "substantially identical" to the GEP polypeptide sequences described herein have an amino acid sequence
5 that is at least 80% (e.g., 85%, 90%, 95%, or 99%) identical to the amino acid sequence of the GEP polypeptides represented by the SEQ ID NOs. listed in Table 1. For purposes of comparison, the length of the reference GEP polypeptide sequence will generally be at least 16 amino acids, e.g., at least 20 or 25 amino acids.

10 In the case of polypeptide sequences that are less than 100% identical to a reference sequence, the non-identical positions are preferably, but not necessarily, conservative substitutions for the reference sequence. Conservative substitutions typically include substitutions within the following groups: glycine and alanine; valine, isoleucine, and leucine; aspartic acid and glutamic acid; asparagine and
15 glutamine; serine and threonine; lysine and arginine; and phenylalanine and tyrosine.

Where a particular polypeptide is said to have a specific percent identity to a reference polypeptide of a defined length, the percent identity is relative to the reference polypeptide. Thus, a polypeptide that is 50% identical to a reference
20 polypeptide that is 100 amino acids long can be a 50 amino acid polypeptide that is completely identical to a 50 amino acid long portion of the reference polypeptide. It also might be a 100 amino acid long polypeptide which is 50% identical to the reference polypeptide over its entire length. Of course, other polypeptides also will meet the same criteria.

25 The invention also features purified or isolated antibodies that specifically bind to a GEP polypeptide. By "specifically binds" is meant that an antibody recognizes and binds to a particular antigen, e.g., a GEP polypeptide, but does not substantially recognize and bind to other molecules in a sample, e.g., a biological sample that naturally includes a GEP polypeptide.

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In another aspect, the invention features a method for detecting a GEP polypeptide in a sample. This method includes: obtaining a sample suspected of containing a GEP polypeptide; contacting the sample with an antibody that specifically binds to a GEP polypeptide under conditions that allow the formation
5 of complexes of an antibody and the GEP polypeptide; and detecting the complexes, if any, as an indication of the presence of a GEP polypeptide in the sample.

Also encompassed by the invention is a method of obtaining a gene related to (i.e., a functional homolog or ortholog of) a GEP gene. Such a method entails
10 obtaining a labeled probe that includes an isolated nucleic acid which encodes all or a portion of a GEP nucleic acid, or a homolog or ortholog thereof; screening a nucleic acid fragment library with the labeled probe under conditions that allow hybridization of the probe to nucleic acid fragments in the library, thereby forming nucleic acid duplexes; isolating labeled duplexes, if any; and preparing a full-length
15 gene sequence from the nucleic acid fragments in any labeled duplex to obtain a gene related to the GEP gene.

The invention offers several advantages. For example, the methods for identifying antibacterial agents can be configured for high throughput screening of numerous candidate antibacterial agents.

20 Unless otherwise defined, all technical and scientific terms used herein have the same meaning as commonly understood by one of ordinary skill in the art to which this invention belongs. Although methods and materials similar or equivalent to those described herein can be used in the practice or testing of the present invention, suitable methods and materials are described herein. All
25 publications, patent applications, patents, and other references mentioned herein are incorporated herein by reference in their entirety. In the case of a conflict, the present specification, including definitions, will control. In addition, the materials, methods, and examples are illustrative and are not intended to limit the scope of the invention, which is defined by the claims.

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Other features and advantages of the invention will be apparent from the following detailed description, and from the claims.

Brief Description of the Drawings

Fig. 1 is a representation of the amino acid and coding strand and non-coding strand nucleic acid sequences of the gep103 polypeptide and gene from a *Streptococcus pneumonia* strain (SEQ ID NOs:1, 2, and 3 respectively).

Fig. 2 is a representation of the amino acid and coding strand and non-coding strand nucleic acid sequences of the gep1119 polypeptide and gene from a *Streptococcus pneumonia* strain (SEQ ID NOs:4, 5 and 6, respectively).

Fig. 3 is a representation of the amino acid and coding strand and non-coding strand nucleic acid sequences of the gep1122 polypeptide and gene from a *Streptococcus pneumonia* strain (SEQ ID NOs:7, 8, and 9, respectively).

Fig. 4 is a representation of the amino acid and coding strand and non-coding strand nucleic acid sequences of the gep1315 polypeptide and gene from a *Streptococcus pneumonia* strain (SEQ ID NOs:10, 11, and 12, respectively).

Fig. 5 is a representation of the amino acid and coding strand and non-coding strand nucleic acid sequences of the gep1493 polypeptide and gene from a *Streptococcus pneumonia* strain (SEQ ID NOs:13, 14, and 15, respectively).

Fig. 6 is a representation of the amino acid and coding strand and non-coding strand nucleic acid sequences of the gep1507 polypeptide and gene from a *Streptococcus pneumonia* (SEQ ID NOs:16, 17, and 18, respectively).

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Fig. 7 is a representation of the amino acid and coding strand and non-coding strand nucleic acid sequences of the gep1511 polypeptide and gene from a *Streptococcus pneumonia* (SEQ ID NOs:19, 20, and 21, respectively).

Fig. 8 is a representation of the amino acid and coding strand and non-coding strand nucleic acid sequences of the gep1518 polypeptide and gene from a *Streptococcus pneumonia* (SEQ ID NOs:22, 23, and 24, respectively).

Fig. 9 is a representation of the amino acid and coding strand and non-coding strand nucleic acid sequences of the gep1546 polypeptide and gene from a *Streptococcus pneumonia* strain (SEQ ID NOs:25, 26, and 27, respectively).

Fig. 10 is a representation of the amino acid and coding strand and non-coding strand nucleic acid sequences of the gep1551 polypeptide and gene from a *Streptococcus pneumonia* strain (SEQ ID NOs:28, 29, and 30, respectively).

Fig. 11 is a representation of the amino acid and coding strand and non-coding strand nucleic acid sequences of the gep1561 polypeptide and gene from a *Streptococcus pneumonia* strain (SEQ ID NOs:31, 32, and 33, respectively).

Fig. 12 is a representation of the amino acid and coding strand and non-coding strand nucleic acid sequences of the gep1580 polypeptide and gene from a *Streptococcus pneumonia* strain (SEQ ID NOs:34, 35, and 36, respectively).

Fig. 13 is a representation of the amino acid and coding strand and non-coding strand nucleic acid sequences of the gep1713 polypeptide and gene from a *Streptococcus pneumonia* (SEQ ID NOs:37, 38, and 39, respectively).

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Fig. 14 is a representation of the amino acid and coding strand and non-coding strand nucleic acid sequences of the gep222 polypeptide and gene from a *Streptococcus pneumonia* (SEQ ID NOs:40, 41, and 42, respectively).

Fig. 15 is a representation of the amino acid and coding strand and non-coding strand nucleic acid sequences of the gep2283 polypeptide and gene from a *Streptococcus pneumonia* (SEQ ID NOs:43, 44, and 45, respectively).

Fig. 16 is a representation of the amino acid and coding strand and non-coding strand nucleic acid sequences of the gep273 polypeptide and gene from a *Streptococcus pneumonia* strain (SEQ ID NOs:46, 47, and 48, respectively).

Fig. 17 is a representation of the amino acid and coding strand and non-coding strand nucleic acid sequences of the gep286 polypeptide and gene from a *Streptococcus pneumonia* strain (SEQ ID NOs:49, 50, and 51, respectively).

Fig. 18 is a representation of the amino acid and coding strand and non-coding strand nucleic acid sequences of the gep311 polypeptide and gene from a *Streptococcus pneumonia* (SEQ ID NOs:52, 53, and 54, respectively).

Fig. 19 is a representation of the amino acid and coding strand and non-coding strand nucleic acid sequences of the gep3262 polypeptide and gene from a *Streptococcus pneumonia* (SEQ ID NOs:55, 56, and 57, respectively).

Fig. 20 is a representation of the amino acid and coding strand and non-coding strand nucleic acid sequences of the gep3387 polypeptide and gene from a *Streptococcus pneumonia* (SEQ ID NOs:58, 59, and 60, respectively).

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Fig. 21 are a representation of the amino acid and coding strand and non-coding strand nucleic acid sequences of the gep47 polypeptide and gene from a *Streptococcus pneumonia* strain (SEQ ID NOs:61, 62, and 63, respectively).

Fig. 22 is a representation of the amino acid and coding strand and non-coding strand nucleic acid sequences of the gep61 polypeptide and gene from a *Streptococcus pneumonia* strain (SEQ ID NOs:64, 65, and 66, respectively).

Fig. 23 is a representation of the amino acid and coding strand and non-coding strand nucleic acid sequences of the gep76 polypeptide and gene from a *Streptococcus pneumonia* strain (SEQ ID NOs:67, 68, and 69, respectively).

Fig. 24 is a representation of the amino acid and coding strand and non-coding strand nucleic acid sequences of the B-yneS polypeptide and gene from a *Bacillus subtilis* strain (SEQ ID NOs:70, 71, and 72, respectively).

Fig. 25 is a schematic representation of the PCR strategy used to produce DNA molecules used for targeted deletions of essential genes in *Streptococcus pneumoniae*.

Fig. 26 is a schematic representation of the strategy used to produce targeted deletions of essential genes in *Streptococcus pneumoniae*.

Detailed Description of the Invention

Identifying *Streptococcus* Genes in Essential Operons

As shown by the experiments described below, each of the GEP genes is located within an operon that is essential for survival of *Streptococcus pneumonia*. *Streptococcus pneumonia* is available from the ATCC. To identify genes located within essential operons, mutants of *Streptococcus pneumonia* were produced. In

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general, mutagenesis of *Streptococcus pneumonia* can be accomplished using any of various art-known methods.

In general, and for the examples set forth below, genes located within essential *Streptococcus pneumonia* operons can be identified using genes from a

5 *Streptococcus pneumonia* RX1 genomic library, which was produced using standard methods (see Kim et al., Nucl. Acids. Res. 20: 1083-1085 (1992) and Ausubel et al. (eds.), 1995, Current Protocols in Molecular Biology, (John Wiley & Sons, NY)). Genes in this *Streptococcus* library were disrupted using a shuttle mutagenesis approach with the transposon TnPho-A. Each disrupted gene then was

10 tested to determine whether it was located within an operon that is essential for survival of *Streptococcus pneumonia*. In this method, 2 ml of LB broth supplemented with chloramphenicol (10 µg/ml), MgSO₄ (10 mM) and maltose (0.2%) were inoculated with 50 µl of the *Streptococcus pneumonia* RX-1 plasmid library. The culture was grown at 37°C while shaking until the OD₆₅₀ of the

15 culture reached 0.8 (approximately 2 hours). A 1 ml aliquot of TnPho-A-containing phage (10⁹ pfu/ml) was added to 1 ml of the *Streptococcus* culture, producing a ratio of approximately 10 phage to 1 cell. The phage and cells were incubated at 37°C for 30 minutes. A 4 ml aliquot of LB broth, warmed to 37°C, then was added to the phage/cell mixture, and the mixture was incubated at 37°C,

20 while shaking, for 1 hour. The cells then were pelleted by centrifuging them at 3500 rpm in a Beckman tabletop centrifuge for 5 minutes.

The pelleted cells then were resuspended in 800 µl of LB broth, and a 200 µl aliquot of cells was plated onto each of four petri plates containing LB agar supplemented with chloramphenicol (10 µg/ml), kanamycin (50 µg/ml), and

25 erythromycin (300 µg/ml). The plates then were incubated overnight at 37°C, and the number of colonies appearing on the plates was counted. Approximately 18,000 colonies then were pooled and used to inoculate 50 ml of LB broth, which was incubated overnight at 37°C. Plasmid DNA from the culture then was extracted using a Qiagen MIDI Prep Kit; other art-known extraction methods can

30 be substituted.

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The concentration of the extracted DNA was measured, and 100 ng of the DNA was transformed, by electroporation, into *E. coli* DH10B cells (Gibco BRL). A 1 ml aliquot of SOC broth then was added the transformed cells, and the cells were incubated at 37°C for 1 hour before being pelleted by centrifugation at 3500
5 RPM for 5 minutes. The cells then were resuspended in 200 µl of LB broth, and aliquots of 2, 20, and 50 µl were plated onto petri plates containing LB agar and antibiotics as described above. After incubating the plates overnight at 37°C, 93 colonies were picked and used, individually, to inoculate 1.25 ml of Terrific broth supplemented with chloramphenicol (10µg/ml), kanamycin (50µg/ml), and
10 erythromycin (300µg/ml). The cultures were incubated at 37°C for approximately 20 hours, while shaking. The DNA from each culture then was extracted, using a conventional alkaline lysis miniprep method.

The extracted DNA samples then were used, individually, to transform *Streptococcus pneumonia* cells in a 96-well microtitre format. The transposon
15 promotes insertion of the mutagenized gene into the bacterial chromosome. Non-transforming clones indicate that the mutation was within an operon containing an essential gene.

The non-transforming clones then were grown in 50 ml of Terrific broth supplemented with chloramphenicol (10 µg/ml), kanamycin (50 µg/ml), and
20 erythromycin (300 µg/ml). DNA from these clones was extracted and retransformed into *Streptococcus pneumonia* and plated on petri dishes to confirm that they were non-transforming. The genes located within essential operons then were sequenced, using primers that hybridize to sequences of the transposon. The sequences of the primers were: 5'GCAGCCCGGTTTCCAGAACAGG3' (SEQ ID
25 NO: 73) and 5'GATTTAGCCCGAGTCGGCCGCACG3' (SEQ ID NO: 74).

In an alternative method, which also was used, the transposon Tn 10 was used to disrupt genes in a *Streptococcus pneumonia* fosmid library, which was produced using standard methods. A 50 ml aliquot of TBMM broth supplemented with chloramphenicol (10µg/ml), MgSO₄ (10 mM), and maltose (0.2%) were
30 inoculated with a single fosmid colony from the fosmid library, and the cultures

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were grown overnight at 37°C. The cells then were pelleted and resuspended in 5 ml of LB broth supplemented with chloramphenicol (10 µg/ml), MgSO₄ (10 mM), and maltose (0.2%). A 100 µl aliquot of the cells then was mixed with 100 µl of Tn10 phage lysate (10¹⁰ pfu/ml), and the mixture was incubated at room

5 temperature for 15 minutes and then incubated at 37°C for 15 minutes.

A 5 ml aliquot of LB broth supplemented with IPTG (1 mM) and sodium citrate (50 mM) and warmed to 37°C then was added to the cell/phage mixture. After incubating the cell/phage mixture at 37°C, while shaking, the cells were pelleted and resuspended in 800 µl of LB broth. The cells then were plated onto 4
10 plates of LB agar supplemented with chloramphenicol (10 µg/ml) and erythromycin (300 µg/ml). After incubating the cells overnight at 37°C, at least 10,000 of the resulting colonies were used to inoculate 50 ml of LB broth. DNA then was extracted and quantified using standard methods, and 100 ng of DNA were used to transform *E. coli* DH10B cells (Gibco BRL) via electroporation. After adding 1 ml
15 of SOC broth to the cells, the cells were incubated at 37°C for 1 hour. The cells then were pelleted and suspended in 200 µl LB broth, and aliquots of 2, 20, and 50 µl were plated onto LB agar supplemented with chloramphenicol (10 µg/ml), kanamycin (50 µg/ml), and erythromycin (300 µg/ml). The plates then were incubated overnight at 37°C, and 93 colonies were picked and used to inoculate
20 1.25 ml of Terrific broth supplemented with chloramphenicol (10 µg/ml), kanamycin (50 µg/ml) and erythromycin (300 µg/ml). These cultures were incubated for approximately 20 hours, while shaking, and the DNA was isolated using a standard miniprep method. The extracted DNA then was used to transform *Streptococcus pneumonia*, and the genes located within essential operons were
25 sequenced as described above. The sequences of the primers used for sequencing were: 5'CCGCCATTCTTTGCTGTTTCG3' (SEQ ID NO: 75) and 5'TTACACGTTACTAAAGGGAATG3' (SEQ ID NO: 76).

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Identification of the gep1493, gep1507, gep1546, gep273, gep286, and gep76 Genes as Essential Genes

As shown by the experiments described below, the gep1493, gep1507, gep1546, gep273, gep286, and gep76 genes each have been shown to be essential for survival of *Streptococcus pneumoniae*. Each of the gep1493, gep1507, gep1546, gep273, gep286, and gep76 genes has been identified as essential by creating a targeted deletion of each gene, separately, in *Streptococcus pneumoniae*.

Each of the gep1493, gep1507, gep1546, gep273, gep286, and gep76 genes was, separately, replaced with a nucleic acid sequence conferring resistance to the antibiotic erythromycin (an "erm" gene). Other genetic markers can be used in lieu of this particular antibiotic resistance marker. Polymerase chain reaction (PCR) amplification was used to make a targeted deletion in the *Streptococcus* genomic DNA, as shown in Fig. 25. Several PCR reactions were used to produce the DNA molecules needed to carry out target deletion of the genes of interest. First, using primers 5 and 6, an erm gene was amplified from pIL252 from *B. subtilis* (available from the *Bacillus* Genetic Stock Center, Columbus, OH). Primer 5 consists of 21 nucleotides that are identical to the promoter region of the erm gene and complementary to Sequence A. Primer 5 has the sequence 5'GTG TTC GTG CTG ACT TGC ACC3' (SEQ ID NO: 77). Primer 6 consists of 21 nucleotides that are complementary to the 3' end of the erm gene. Primer 6 has the sequence 5'GAA TTA TTT CCT CCC GTT AAA3' (SEQ ID NO: 78). PCR amplification of the erm gene was carried out under the following conditions: 30 cycles of 94°C for 1 minute, 55°C for 1 minute, and 72°C for 1.5 minutes, followed by one cycle of 72°C for 10 minutes.

In the second and third PCR reactions, sequences flanking the gene of interest were amplified and produced as hybrid DNA molecules that also contained a portion of the erm gene. The second reaction produced a double-stranded DNA molecule (termed "Left Flanking Molecule") that includes sequences upstream of the 5' end of the gene of interest and the first 21 nucleotides of the erm gene. As shown in Fig. 25, this reaction utilized primer 1, which is 21 nucleotides in length

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and identical to a sequence that is located approximately 500 bp upstream of the translation start site of the gene of interest. Primers 1 and 2 are gene-specific and include the sequences 5'CTC CGT GAA GTC CAC CTG AT3' (SEQ ID NO:79) and 5'GGT GCA AGT CAG CAC GAA CAC GCG ACA TAG GTT CCA GTT
5 AGG3' (SEQ ID NO:80), respectively, for gep1493. Primer 2 is 42 nucleotides in length, with 21 of the nucleotides at the 3' end of the primer being complementary to the 5' end of the sense strand of the gene of interest. The 21 nucleotides at the 5' end of the primer were identical to Sequence A and are therefore complementary to the 5' end of the *erm* gene. Thus, PCR amplification using primers 1 and 2
10 produced the left flanking DNA molecule, which is a hybrid DNA molecule containing a sequence located upstream of the gene of interest and 21 base pairs of the *erm* gene, as shown in Fig. 25.

The third PCR reaction was similar to the second reaction, but produced the right flanking DNA molecule, shown in Fig. 25. The right flanking DNA molecule
15 contains 21 base pairs of the 3' end of the *erm* gene, a 21 base pair portion of the 3' end of the gene of interest, and sequences downstream of the gene of interest. This right flanking DNA molecule was produced with gene-specific primers 3 and 4. For gep 1493, primers 3 and 4 included the sequences 5'TTT AAC GGG AGG AAA TAA TTC CCA TAT CGT GGC TCC TGA AT 3' (SEQ ID NO:81) and
20 5'TAA AGC CCT CAT GTC GAA CC3' (SEQ ID NO:82), respectively. Primer 3 is 42 nucleotides; the 21 nucleotides at the 5' end of Primer 3 are identical to Sequence B and therefore are identical to the 3' end of the *erm* gene. The 21 nucleotides at the 3' end of Primer 3 are identical to the 3' end of the gene of interest. Primer 4 is 21 nucleotides in length and is complementary to a sequence
25 located approximately 500 bp downstream of the gene of interest. As discussed above, primers 1-4 are gene-specific, and the sequences disclosed above were used for gep1493. Gene-specific primers were used to identify the other essential genes described herein, as shown in Table 2.

TABLE 2: Primers Used in Identifying Essential Genes

Gene	Primer 1	Primer 2	Primer 3	Primer 4
gepl493	5'CTCCGTGAA GTCCACCTGA T3' (SEQ ID NO:79)	5'GGTGCAAGT CAGCACGAAC ACTGCTCGCG TAGATTGATT TG3' (SEQ ID NO:80)	5'TTTAACGGG AGGAAATAAT TCGGGGATTG AACCTAACCC AT3' (SEQ ID NO:81)	5'TTGGCAAG AAGGCAGAG AAT3' (SEQ ID NO:82)
gepl507	5'GCATGAGAA ACCCAGTCTC C3' (SEQ ID NO:83)	5'GGTGCAAGT CAGCACGAAC ACGCGACATA GGTTCCAGTT AGG3' (SEQ ID NO:84)	5'TTTAACGGG AGGAAATAAT TCCCATATCG TGGCTCCTGA AT3' (SEQ ID NO:85)	5'TAAAGCCC TCATGTCGAA CC3' (SEQ ID NO:86)
gepl546	5'CAGTGACGA TACAGATGAA GAA3' (SEQ ID NO:87)	5'GGTGCAAGT CAGCACGAAC ACGATGCTGG CTTCGTTGAG TG3' (SEQ ID NO:88)	5'TTTAACGGG AGGAAATAAT TCGTCGCGAC TCCTAGCCAT AC3' (SEQ ID NO:89)	5'CCAGCAAA GGAAAACCG ATA3' (SEQ ID NO:90)
gep273	5'GGTCAGTGA CAGCAGCAGA T3' (SEQ ID NO:91)	5'GGTGCAAGT CAGCACGAAC ACGGCCTTGG AAAAAAGACC AT3' (SEQ ID NO:92)	5'TTTAACGGG AGGAAATAAT TCCCGCTTAA ATTCTGCCAA TC3' (SEQ ID NO:93)	5'CCCATAAC CGTATCACCT GG3' (SEQ ID NO:94)
gep286	5'CGGAACGGC TATGAAAAAA A3' (SEQ ID NO:95)	5'GGTGCAAGT CAGCACGAAC ACACGACGAA AGGCAACCAT AC3' (SEQ ID NO:96)	5'TTTAACGGG AGGAAATAAT TCTGGTATGG GGGTTGATGA AG3' (SEQ ID NO:97)	5'TCGCCCTAC TTTTCGTATG C3' (SEQ ID NO:98)
gep76	5'AGCGATATT AGTGC GGGAG A3' (SEQ ID NO:99)	5'GGTGCAAGT CAGCACGAAC ACCAGCAATT TTGTCATCAG TCG3' (SEQ ID NO:100)	5'TTTAACGGG AGGAAATAAT TCCTGGGGTA ATGGAGCACA GT3' (SEQ ID NO:101)	5'GGGATTGT CACGGTAAA ACC3' (SEQ ID NO:102)

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PCR amplification of the left and right flanking DNA molecules was carried out, separately, in 50 μ l reaction mixtures containing: 1 μ l *Streptococcus pneumoniae* (RX1) DNA (0.25 μ g), 2.5 μ l Primer 1 or Primer 4 (10 pmol/ μ l), 2.5 μ l Primer 2 or Primer 3 (20 pmol/ μ l), 1.2 μ l a mixture dNTPS (10 mM each), 5 37 μ l H₂O, 0.7 μ l Taq polymerase (5 U/ μ l), and 5 μ l 10x Taq polymerase buffer (10 mM Tris, 50 mM KCl, 2.5 mM MgCl₂). The left and right flanking DNA molecules were amplified using the following PCR cycling program: 95°C for 2 minutes; 72°C for 1 minute; 94°C for 30 seconds; 49°C for 30 seconds; 72°C for 1 minute; repeating the 94°C, 49°C, and 72°C incubations 30 times; 72°C for 10 10 minutes and then stopping the reactions. A 15 μ l aliquot of each reaction mixture then was electrophoresed through a 1.2% low melting point agarose gel in TAE buffer and then stained with ethidium bromide. Fragments containing the amplified left and right flanking DNA molecules were excised from the gel and purified using the QIAQUICK™ gel extraction kit (Qiagen, Inc.) Other art-known methods 15 for amplifying and isolating DNA can be substituted. The flanking left and right DNA fragments were eluted into 30 μ l TE buffer at pH 8.0.

The amplified *erm* gene and left and right flanking DNA molecules were then fused together to produce the fusion product, as shown in Fig. 25. The fusion PCR reaction was carried out in a volume of 50 μ l containing: 2 μ l of each of the 20 left and right flanking DNA molecules and the *erm* gene PCR product; 5 μ l of 10x buffer; 2.5 μ l of Primer 1 (10 pmol/ μ l); 2.5 μ l of Primer 4 (10 pmol/ μ l), 1.2 μ l dNTP mix (10 mM each) 32 μ l H₂O, and 0.7 μ l Taq polymerase. The PCR reaction was carried out using the following cycling program: 95°C for 2 minutes; 72°C for 1 minute; 94°C for 30 seconds, 48°C for 30 seconds; 72°C for 3 minutes; 25 repeat the 94°C, 48°C and 72°C incubations 25 times; 72°C for 10 minutes. After the reaction was stopped, a 12 μ l aliquot of the reaction mixture was electrophoresed through an agarose gel to confirm the presence of a final product of approximately 2 kb.

A 5 μ l aliquot of the fusion product was used to transform *S. pneumoniae* 30 grown on a medium containing erythromycin in accordance with standard

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techniques. As shown in Fig. 26, the fusion product and the *S. pneumoniae* genome undergo a homologous recombination event so that the *erm* gene replaces the chromosomal copy of the gene of interest, thereby creating a gene knockout. Disruption of an essential gene results in no growth on a medium containing
5 erythromycin. Using this gene knockout method, the gep1493, gep1507, gep1546, gep273, gep286, and gep76 genes were each identified as being essential for survival.

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Identification of Homologs and Orthologs of GEP Polypeptides

Having shown that the various GEP genes are essential or located within operons that are essential for survival of *Streptococcus*, it can be expected that homologs and orthologs of the polypeptides encoded by these genes, when present

5 in other organisms, for example *B. subtilis*, are essential or located within operons that are essential for survival of that organism as well, and therefore are useful targets for identifying antibacterial agents. Using the sequences of the GEP polypeptides identified in *Streptococcus*, homologs and orthologs of these polypeptides can be identified in other organisms. For example, the coding

10 sequences of the GEP nucleic acids can be used to search the GenBank database of nucleotide sequences to identify homologs or orthologs that are expressed from essential operons in other organisms. Sequence comparisons can be performed using the Basic Local Alignment Search Tool (BLAST) (Altschul et al., *J. Mol. Biol.*, 215:403-410 1990). The percent sequence identity shared by the GEP

15 polypeptides and their homologs or orthologs can be determined using the GAP program from the Genetics Computer Group (GCG) Wisconsin Sequence Analysis Package (Wisconsin Package Version 9.0, GCG; Madison, WI). The following parameters are suitable: gap creation penalty, 12 (protein) 50 (DNA); gap extension penalty, 4 (protein) 3 (DNA). Typically, the GEP polypeptides and their

20 homologs share at least 25% (e.g., at least 40%) sequence identity. Typically, the DNA sequences encoding GEP polypeptides and their homologs share at least 35% (e.g., at least 45%) sequence identity. To confirm that the homologs or orthologs of the GEP polypeptides are expressed from operons that are essential for survival of bacteria, the operon encoding each of the homologs or orthologs can be,

25 separately, deleted from the genome of the host organism.

Identification of Essential Operons in Additional *Streptococcus* Strains

Now that the various GEP genes have been identified as being located within operons that are essential for survival, these genes, or fragments thereof, can be used to detect homologous or orthologous genes in other organisms. In

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particular, these genes can be used to analyze various pathogenic and non-pathogenic strains of bacteria. Fragments of a nucleic acid (DNA or RNA) encoding a GEP polypeptide or homolog or ortholog (or sequences complementary thereto) can be used as probes in conventional nucleic acid hybridization assays of pathogenic bacteria. For example, nucleic acid probes (which typically are 8-30, or usually 15-20, nucleotides in length) can be used to detect GEP genes or homologs or orthologs thereof in art-known molecular biology methods, such as Southern blotting, Northern blotting, dot or slot blotting, PCR amplification methods, colony hybridization methods, and the like. Typically, an oligonucleotide probe based on the nucleic acid sequences described herein, or fragments thereof, is labeled and used to screen a genomic library constructed from mRNA obtained from a *Streptococcus* or bacterial strain of interest. A suitable method of labeling involves using polynucleotide kinase to add ^{32}P -labeled ATP to the oligonucleotide used as the probe. This method is well known in the art, as are several other suitable methods (e.g., biotinylation and enzyme labeling).

Hybridization of the oligonucleotide probe to the library, or other nucleic acid sample, typically is performed under stringent to highly stringent conditions. Nucleic acid duplex or hybrid stability is expressed as the melting temperature or T_m , which is the temperature at which a probe dissociates from a target DNA. This melting temperature is used to define the required stringency conditions. If sequences are to be identified that are related and substantially identical to the probe, rather than identical, then it is useful to first establish the lowest temperature at which only homologous hybridization occurs with a particular concentration of salt (e.g., SSC or SSPE). Then, assuming that 1% mismatching results in a 1°C decrease in the T_m , the temperature of the final wash in the hybridization reaction is reduced accordingly (for example, if sequences having $\geq 95\%$ identity with the probe are sought, the final wash temperature is decreased by 5°C). In practice, the change in T_m can be between 0.5° and 1.5°C per 1% mismatch.

As used herein, highly stringent conditions refer to hybridization at 68°C in 5x SSC/5x Denhardt's solution/1.0% SDS, and washing in 0.2x SSC/0.1% SDS at

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42°C. Stringent conditions refer to washing in 3x SSC at 42°C. The parameters of salt concentration and temperature can be varied to achieve the optimal level of identity between the probe and the target nucleic acid. Additional guidance regarding such conditions is readily available in the art, for example, by Sambrook et al., 1989, *Molecular Cloning, A Laboratory Manual*, Cold Spring Harbor Press, N.Y.; and Ausubel et al. (eds.), 1995, *Current Protocols in Molecular Biology*, (John Wiley & Sons, N.Y.) at Unit 2.10.

In one approach, libraries constructed from pathogenic or non-pathogenic *Streptococcus* or bacterial strains can be screened. For example, such strains can be screened for expression of GEP genes by Northern blot analysis. Upon detection of transcripts of the GEP genes or homologs or orthologs thereof, libraries can be constructed from RNA isolated from the appropriate strain, utilizing standard techniques well known to those of skill in the art. Alternatively, a total genomic DNA library can be screened using an GEP gene probe (or a probe directed to a homolog or ortholog thereof).

New gene sequences can be isolated, for example, by performing PCR using two degenerate oligonucleotide primer pools designed on the basis of nucleotide sequences within the GEP genes, or their homologs or orthologs, as depicted herein. The template for the reaction can be DNA obtained from strains known or suspected to express a GEP allele or an allele of a homolog or ortholog thereof. The PCR product can be subcloned and sequenced to ensure that the amplified sequences represent the sequences of a new GEP nucleic acid sequence, or a sequence of a homolog or ortholog thereof.

Synthesis of the various GEP polypeptides or their homologs or orthologs (or an antigenic fragment thereof) for use as antigens, or for other purposes, can readily be accomplished using any of the various art-known techniques. For example, a polypeptide or homolog or ortholog thereof, or an antigenic fragment(s), can be synthesized chemically *in vitro*, or enzymatically (e.g., by *in vitro* transcription and translation). Alternatively, the gene can be expressed in, and the polypeptide purified from, a cell (e.g., a cultured cell) by using any of the

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numerous, available gene expression systems. For example, the polypeptide antigen can be produced in a prokaryotic host (e.g., *E. coli* or *B. subtilis*) or in eukaryotic cells, such as yeast cells or insect cells (e.g., by using a baculovirus-based expression vector).

- 5 Proteins and polypeptides can also be produced in plant cells, if desired. For plant cells viral expression vectors (e.g., cauliflower mosaic virus and tobacco mosaic virus) and plasmid expression vectors (e.g., Ti plasmid) are suitable. Such cells are available from a wide range of sources (e.g., the American Type Culture Collection, Rockland, MD; also, *see*, e.g., Ausubel et al., *Current Protocols in*
- 10 *Molecular Biology*, John Wiley & Sons, New York, 1994). The optimal methods of transformation or transfection and the choice of expression vehicle will depend on the host system selected. Transformation and transfection methods are described, e.g., in Ausubel et al., *supra*; expression vehicles may be chosen from those provided, e.g., in *Cloning Vectors: A Laboratory Manual* (P.H. Pouwels et
- 15 al., 1985, Supp. 1987). The host cells harboring the expression vehicle can be cultured in conventional nutrient media, adapted as needed for activation of a chosen gene, repression of a chosen gene, selection of transformants, or amplification of a chosen gene.

- If desired, GEP polypeptides or their homologs or orthologs can be
- 20 produced as fusion proteins. For example, the expression vector pUR278 (Ruther et al., *EMBO J.*, 2:1791, 1983) can be used to create *lacZ* fusion proteins. The art-known pGEX vectors can be used to express foreign polypeptides as fusion proteins with glutathione S-transferase (GST). In general, such fusion proteins are soluble and can be easily purified from lysed cells by adsorption to glutathione-
- 25 agarose beads followed by elution in the presence of free glutathione. The pGEX vectors are designed to include thrombin or factor Xa protease cleavage sites so that the cloned target gene product can be released from the GST moiety.

- In an exemplary insect cell expression system, a baculovirus such as *Autographa californica* nuclear polyhedrosis virus (AcNPV), which grows in
- 30 *Spodoptera frugiperda* cells, can be used as a vector to express foreign genes. A

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coding sequence encoding a GEP polypeptide or homolog or ortholog can be cloned into a non-essential region (for example the polyhedrin gene) of the viral genome and placed under control of a promoter, e.g., the polyhedrin promoter or an exogenous promoter. Successful insertion of a gene encoding a GEP

5 polypeptide or homolog or ortholog can result in inactivation of the polyhedrin gene and production of non-occluded recombinant virus (i.e., virus lacking the proteinaceous coat encoded by the polyhedrin gene). These recombinant viruses are then used to infect insect cells (e.g., *Spodoptera frugiperda* cells) in which the inserted gene is expressed (see, e.g., Smith et al., *J. Virol.*, 46:584, 1983; Smith,

10 U.S. Patent No. 4,215,051).

In mammalian host cells, a number of viral-based expression systems can be utilized. When an adenovirus is used as an expression vector, the nucleic acid sequence encoding the GEP polypeptide or homolog or ortholog can be ligated to an adenovirus transcription/ translation control complex, e.g., the late promoter and

15 tripartite leader sequence. This chimeric gene can then be inserted into the adenovirus genome by *in vitro* or *in vivo* recombination. Insertion into a non-essential region of the viral genome (e.g., region E1 or E3) will result in a recombinant virus that is viable and capable of expressing a essential gene product in infected hosts (see, e.g., Logan, Proc. Natl. Acad. Sci. USA, 81:3655, 1984).

20 Specific initiation signals may be required for efficient translation of inserted nucleic acid sequences. These signals include the ATG initiation codon and adjacent sequences. In general, exogenous translational control signals, including, perhaps, the ATG initiation codon, should be provided. Furthermore, the initiation codon must be in phase with the reading frame of the desired coding

25 sequence to ensure translation of the entire sequence. These exogenous translational control signals and initiation codons can be of a variety of origins, both natural and synthetic. The efficiency of expression may be enhanced by the inclusion of appropriate transcription enhancer elements, or transcription terminators (Bittner et al., *Methods in Enzymol.*, 153:516, 1987).

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The GEP polypeptides and homologs and orthologs can be expressed individually or as fusions with a heterologous polypeptide, such as a signal sequence or other polypeptide having a specific cleavage site at the N-and/or C-terminus of the protein or polypeptide. The heterologous signal sequence selected
5 should be one that is recognized and processed, i.e., cleaved by a signal peptidase, by the host cell in which the fusion protein is expressed.

A host cell can be chosen that modulates the expression of the inserted sequences, or modifies and processes the gene product in a specific, desired fashion. Such modifications and processing (e.g., cleavage) of protein products
10 may facilitate optimal functioning of the protein. Various host cells have characteristic and specific mechanisms for post-translational processing and modification of proteins and gene products. Appropriate cell lines or host systems familiar to those of skill in the art of molecular biology can be chosen to ensure the correct modification and processing of the foreign protein expressed. To this
15 end, eukaryotic host cells that possess the cellular machinery for proper processing of the primary transcript, and phosphorylation of the gene product can be used. Such mammalian host cells include, but are not limited to, CHO, VERO, BHK, HeLa, COS, MDCK, 293, 3T3, WI38, and choroid plexus cell lines.

If desired, the GEP polypeptide or homolog or ortholog thereof can be
20 produced by a stably-transfected mammalian cell line. A number of vectors suitable for stable transection of mammalian cells are available to the public, *see*, e.g., Pouwels et al. (supra); methods for constructing such cell lines are also publicly known, e.g., in Ausubel et al. (supra). In one example, DNA encoding the protein is cloned into an expression vector that includes the dihydrofolate reductase
25 (DHFR) gene. Integration of the plasmid and, therefore, the GEP polypeptide-encoding gene into the host cell chromosome is selected for by including 0.01-300 μ M methotrexate in the cell culture medium (as described in Ausubel et al., supra). This dominant selection can be accomplished in most cell types.

Recombinant protein expression can be increased by DHFR-mediated
30 amplification of the transfected gene. Methods for selecting cell lines bearing gene

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amplifications are described in Ausubel et al. (supra); such methods generally involve extended culture in medium containing gradually increasing levels of methotrexate. DHFR-containing expression vectors commonly used for this purpose include pCVSEII-DHFR and pAdD26SV(A) (described in Ausubel et al.,
5 supra).

A number of other selection systems can be used, including but not limited to, herpes simplex virus thymidine kinase genes, hypoxanthine-guanine phosphoribosyl-transferase genes, and adenine phosphoribosyltransferase genes, which can be employed in *tk*, *hgpri*, or *aprt* cells, respectively. In addition, *gpt*,
10 which confers resistance to mycophenolic acid (Mulligan et al., *Proc. Natl. Acad. Sci. USA*, 78:2072, 1981); *neo*, which confers resistance to the aminoglycoside G-418 (Colberre-Garapin et al., *J. Mol. Biol.*, 150:1, 1981); and *hygro*, which confers resistance to hygromycin (Santerre et al., *Gene*, 30:147, 1981), can be used.

Alternatively, any fusion protein can be readily purified by utilizing an
15 antibody or other molecule that specifically binds to the fusion protein being expressed. For example, a system described in Janknecht et al., *Proc. Natl. Acad. Sci. USA*, 88:8972 (1981), allows for the ready purification of non-denatured fusion proteins expressed in human cell lines. In this system, the gene of interest is subcloned into a vaccinia recombination plasmid such that the gene's open reading
20 frame is translationally fused to an amino-terminal tag consisting of six histidine residues. Extracts from cells infected with recombinant vaccinia virus are loaded onto Ni²⁺ nitriloacetic acid-agarose columns, and histidine-tagged proteins are selectively eluted with imidazole-containing buffers.

Alternatively, a GEP polypeptide or homolog or ortholog, or a portion
25 thereof, can be fused to an immunoglobulin Fc domain. Such a fusion protein can be readily purified using a protein A column, for example. Moreover, such fusion proteins permit the production of a chimeric form of a GEP polypeptide or homolog or ortholog having increased stability *in vivo*.

Once the recombinant GEP polypeptide (or homolog or ortholog) is
30 expressed, it can be isolated (i.e., purified). Secreted forms of the polypeptides can

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be isolated from cell culture media, while non-secreted forms must be isolated from the host cells. Polypeptides can be isolated by affinity chromatography. For example, an anti-gep103 antibody (e.g., produced as described herein) can be attached to a column and used to isolate the protein. Lysis and fractionation of

5 cells harboring the protein prior to affinity chromatography can be performed by standard methods (*see, e.g., Ausubel et al., supra*). Alternatively, a fusion protein can be constructed and used to isolate a GEP polypeptide (e.g., a gep103-maltose binding fusion protein, a gep-103- β -galactosidase fusion protein, or a gep103-trpE fusion protein; *see, e.g., Ausubel et al., supra*; New England Biolabs Catalog,

10 Beverly, MA). The recombinant protein can, if desired, be further purified, e.g., by high performance liquid chromatography using standard techniques (*see, e.g., Fisher, Laboratory Techniques In Biochemistry And Molecular Biology, eds., Work and Burdon, Elsevier, 1980*).

Given the amino acid sequences described herein, polypeptides useful in

15 practicing the invention, particularly fragments of GEP polypeptides can be produced by standard chemical synthesis (e.g., by the methods described in *Solid Phase Peptide Synthesis*, 2nd ed., The Pierce Chemical Co., Rockford, IL, 1984) and used as antigens, for example.

Antibodies

20 The GEP polypeptides (or antigenic fragments or analogs of such polypeptides) can be used to raise antibodies useful in the invention, and such polypeptides can be produced by recombinant or peptide synthetic techniques (*see, e.g., Solid Phase Peptide Synthesis, supra*; Ausubel et al., *supra*). Likewise, antibodies can be raised against the GEP homologs and orthologs. In general, the

25 polypeptides can be coupled to a carrier protein, such as KLH, as described in Ausubel et al., *supra*, mixed with an adjuvant, and injected into a host mammal. Antibodies can be purified, for example, by affinity chromatography methods in which the polypeptide antigen is immobilized on a resin.

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In particular, various host animals can be immunized by injection of a polypeptide of interest. Examples of suitable host animals include rabbits, mice, guinea pigs, and rats. Various adjuvants can be used to increase the immunological response, depending on the host species, including but not limited to Freund's
5 (complete and incomplete adjuvant), adjuvant mineral gels such as aluminum hydroxide, surface active substances such as lysolecithin, pluronic polyols, polyanions, peptides, oil emulsions, keyhole limpet hemocyanin, dinitrophenol, BCG (bacille Calmette-Guerin) and *Corynebacterium parvum*. Polyclonal antibodies are heterogeneous populations of antibody molecules derived from the
10 sera of the immunized animals.

Antibodies useful in the invention include monoclonal antibodies, polyclonal antibodies, humanized or chimeric antibodies, single chain antibodies, Fab fragments, F(ab')₂ fragments, and molecules produced using a Fab expression library.

15 Monoclonal antibodies (mAbs), which are homogeneous populations of antibodies to a particular antigen, can be prepared using the GEP polypeptides or homologs or orthologs thereof and standard hybridoma technology (see, e.g., Kohler et al., *Nature*, 256:495, 1975; Kohler et al., *Eur. J. Immunol.*, 6:511, 1976; Kohler et al., *Eur. J. Immunol.*, 6:292, 1976; Hammerling et al., In Monoclonal
20 Antibodies and T Cell Hybridomas, Elsevier, NY, 1981; Ausubel et al., supra).

In particular, monoclonal antibodies can be obtained by any technique that provides for the production of antibody molecules by continuous cell lines in culture, such as those described in Kohler et al., *Nature*, 256:495, 1975, and U.S. Patent No. 4,376,110; the human B-cell hybridoma technique (Kosbor et al.,
25 *Immunology Today*, 4:72, 1983; Cole et al., *Proc. Natl. Acad. Sci. USA*, 80:2026, 1983); and the EBV-hybridoma technique (Cole et al., Monoclonal Antibodies and Cancer Therapy, Alan R. Liss, Inc., pp. 77-96, 1983). Such antibodies can be of any immunoglobulin class including IgG, IgM, IgE, IgA, IgD, and any subclass thereof. The hybridomas producing the mAbs of this invention can be cultivated in
30 *vitro* or *in vivo*.

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Once produced, polyclonal or monoclonal antibodies are tested for specific recognition of a GEP polypeptide or homolog or ortholog thereof in an immunoassay, such as a Western blot or immunoprecipitation analysis using standard techniques, e.g., as described in Ausubel et al., supra. Antibodies that
5 specifically bind to the GEP polypeptides, or conservative variants and homologs or orthologs thereof, are useful in the invention. For example, such antibodies can be used in an immunoassay to detect a GEP polypeptide in pathogenic or non-pathogenic strains of bacteria.

Preferably, antibodies of the invention are produced using fragments of the
10 GEP polypeptides that appear likely to be antigenic, by criteria such as high frequency of charged residues. In one specific example, such fragments are generated by standard techniques of PCR, and are then cloned into the pGEX expression vector (Ausubel et al., supra). Fusion proteins are expressed in *E. coli* and purified using a glutathione agarose affinity matrix as described in Ausubel, et
15 al., supra.

If desired, several (e.g., two or three) fusions can be generated for each protein, and each fusion can be injected into at least two rabbits. Antisera can be raised by injections in a series, typically including at least three booster injections. Typically, the antisera is checked for its ability to immunoprecipitate a recombinant
20 GEP polypeptide or homolog or ortholog, or unrelated control proteins, such as glucocorticoid receptor, chloramphenicol acetyltransferase, or luciferase.

Techniques developed for the production of "chimeric antibodies" (Morrison et al., *Proc. Natl. Acad. Sci.*, **81**:6851, 1984; Neuberger et al., *Nature*, **312**:604, 1984; Takeda et al., *Nature*, **314**:452, 1984) can be used to splice the genes from a
25 mouse antibody molecule of appropriate antigen specificity together with genes from a human antibody molecule of appropriate biological activity. A chimeric antibody is a molecule in which different portions are derived from different animal species, such as those having a variable region derived from a murine mAb and a human immunoglobulin constant region.

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Alternatively, techniques described for the production of single chain antibodies (U.S. Patent 4,946,778; and U.S. Patents 4,946,778 and 4,704,692) can be adapted to produce single chain antibodies against a GEP polypeptide or homolog or ortholog. Single chain antibodies are formed by linking the heavy and
5 light chain fragments of the Fv region via an amino acid bridge, resulting in a single chain polypeptide.

Antibody fragments that recognize and bind to specific epitopes can be generated by known techniques. For example, such fragments can include but are not limited to F(ab')₂ fragments, which can be produced by pepsin digestion of the
10 antibody molecule, and Fab fragments, which can be generated by reducing the disulfide bridges of F(ab')₂ fragments. Alternatively, Fab expression libraries can be constructed (Huse et al., *Science*, 246:1275, 1989) to allow rapid and easy identification of monoclonal Fab fragments with the desired specificity.

Polyclonal and monoclonal antibodies that specifically bind to GEP
15 polypeptides or homologs or orthologs can be used, for example, to detect expression of a GEP gene or homolog or ortholog in another strain of bacteria. For example, a GEP polypeptide can be readily detected in conventional immunoassays of bacteria cells or extracts. Examples of suitable assays include, without limitation, Western blotting, ELISAs, radioimmune assays, and the like.

20 Assay for Antibacterial Agents

The invention provides a method for identifying an antibacterial agent(s). Although the inventors are not bound by any particular theory as to the biological mechanism involved, the new antibacterial agents are thought to inhibit specifically
(1) the function of a polypeptide(s) encoded by a nucleic acid located within an
25 operon containing a GEP gene, or (2) expression of the a gene located within an operon containing a GEP gene, or homologs or orthologs thereof. Screening for antibacterial agents can be rapidly accomplished by identifying those compounds (e.g., polypeptides or small molecules) that specifically bind to a polypeptide encoded by a nucleic acid located within an operon containing a GEP gene. A

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homolog or ortholog of a GEP polypeptide can be substituted for the GEP polypeptide in the methods summarized herein. Specific binding of a test compound to a polypeptide can be detected, for example, *in vitro* by reversibly or irreversibly immobilizing the test compound(s) on a substrate, e.g., the surface of a well of a 96-well polystyrene microtitre plate. Methods for immobilizing polypeptides and other small molecules are well known in the art. For example, the microtitre plates can be coated with a polypeptide encoded by a nucleic acid located within an operon containing a GEP gene (e.g., a GEP polypeptide or a combination of GEP polypeptides and/or homologs and/or orthologs) by adding the polypeptide(s) in a solution (typically, at a concentration of 0.05 to 1 mg/ml in a volume of 1-100 μ l) to each well, and incubating the plates at room temperature to 37°C for 0.1 to 36 hours. Polypeptides that are not bound to the plate can be removed by shaking the excess solution from the plate, and then washing the plate (once or repeatedly) with water or a buffer. Typically, the polypeptide, homolog, or ortholog is contained in water or a buffer. The plate is then washed with a buffer that lacks the bound polypeptide. To block the free protein-binding sites on the plates, the plates are blocked with a protein that is unrelated to the bound polypeptide. For example, 300 μ l of bovine serum albumin (BSA) at a concentration of 2 mg/ml in Tris-HCl is suitable. Suitable substrates include those substrates that contain a defined cross-linking chemistry (e.g., plastic substrates, such as polystyrene, styrene, or polypropylene substrates from Corning Costar Corp. (Cambridge, MA), for example). If desired, a beaded particle, e.g., beaded agarose or beaded sepharose, can be used as the substrate.

Binding of the test compound to the new polypeptides (or homologs or orthologs thereof) can be detected by any of a variety of art-known methods. For example, an antibody that specifically binds to a GEP polypeptide can be used in an immunoassay. If desired, the antibody can be labeled (e.g., fluorescently or with a radioisotope) and detected directly (*see, e.g., West and McMahon, J. Cell Biol. 74:264, 1977*). Alternatively, a second antibody can be used for detection (e.g., a labeled antibody that binds to the Fc portion of an anti-GEP103 antibody).

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In an alternative detection method, the GEP polypeptide is labeled, and the label is detected (e.g., by labeling a GEP polypeptide with a radioisotope, fluorophore, chromophore, or the like). In still another method, the GEP polypeptide is produced as a fusion protein with a protein that can be detected optically, e.g.,
5 green fluorescent protein (which can be detected under UV light). In an alternative method, the polypeptide (e.g., gep103) can be produced as a fusion protein with an enzyme having a detectable enzymatic activity, such as horse radish peroxidase, alkaline phosphatase, β -galactosidase, or glucose oxidase. Genes encoding all of these enzymes have been cloned and are readily available for use by those of skill
10 in the art. If desired, the fusion protein can include an antigen, and such an antigen can be detected and measured with a polyclonal or monoclonal antibody using conventional methods. Suitable antigens include enzymes (e.g., horse radish peroxidase, alkaline phosphatase, and β -galactosidase), and non-enzymatic polypeptides (e.g., serum proteins, such as BSA and globulins, and milk proteins,
15 such as caseins).

In various *in vivo* methods for identifying polypeptides that bind to GEP polypeptides, the conventional two-hybrid assays of protein/protein interactions can be used (see e.g., Chien et al., *Proc. Natl. Acad. Sci. USA*, **88**:9578, 1991; Fields et al., U.S. Pat. No. 5,283,173; Fields and Song, *Nature*, **340**:245, 1989; Le Douarin
20 et al., *Nucleic Acids Research*, **23**:876, 1995; Vidal et al., *Proc. Natl. Acad. Sci. USA*, **93**:10315-10320, 1996; and White, *Proc. Natl. Acad. Sci. USA*, **93**:10001-10003, 1996). Kits for practicing various two-hybrid methods are commercially available (e.g., from Clontech; Palo Alto, CA).

Generally, the two-hybrid methods involve *in vivo* reconstitution of two
25 separable domains of a transcription factor. The DNA binding domain (DB) of the transcription factor is required for recognition of a chosen promoter. The activation domain (AD) is required for contacting other components of the host cell's transcriptional machinery. The transcription factor is reconstituted through the use of hybrid proteins. One hybrid is composed of the AD and a first protein

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of interest. The second hybrid is composed of the DB and a second protein of interest.

Useful reporter genes are those that are operably linked to a promoter which is specifically recognized by the DB. Typically, the two-hybrid system employs
5 the yeast *Saccharomyces cerevisiae* and reporter genes, the expression of which can be selected under appropriate conditions. Other eukaryotic cells, including mammalian and insect cells, can be used, if desired. The two-hybrid system provides a convenient method for cloning a gene encoding a polypeptide (i.e., a candidate antibacterial agent) that binds to a second, preselected polypeptide (e.g.,
10 gep103). Typically, though not necessarily, a DNA library is constructed such that randomly generated sequences are fused to the AD, and the protein of interest (e.g., gep103) is fused to the DB.

In such two-hybrid methods, two fusion proteins are produced. One fusion protein contains the GEP polypeptide (or homolog or ortholog thereof) fused to
15 either a transactivator domain or DNA binding domain of a transcription factor (e.g., of Gal4). The other fusion protein contains a test polypeptide fused to either the DNA binding domain or a transactivator domain of a transcription factor. Once brought together in a single cell (e.g., a yeast cell or mammalian cell), one of the fusion proteins contains the transactivator domain and the other fusion protein
20 contains the DNA binding domain. Therefore, binding of the GEP polypeptide to the test polypeptide (i.e., candidate antibacterial agent) reconstitutes the transcription factor. Reconstitution of the transcription factor can be detected by detecting expression of a gene (i.e., a reporter gene) that is operably linked to a DNA sequence that is bound by the DNA binding domain of the transcription
25 factor.

The methods described above can be used for high throughput screening of numerous test compounds to identify candidate antibacterial (or anti-bacterial) agents. Having identified a test compound as a candidate antibacterial agent, the candidate antibacterial agent can be further tested for inhibition of bacterial growth
30 *in vitro* or *in vivo* (e.g., using an animal, e.g., rodent, model system) if desired.

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Using other, art-known variations of such methods, one can test the ability of a nucleic acid (e.g., DNA or RNA) used as the test compound to bind to a polypeptide encoded by a nucleic acid sequence located within an operon containing a GEP gene or homolog or ortholog thereof.

- 5 *In vitro*, further testing can be accomplished by means known to those in the art such as an enzyme inhibition assay or a whole-cell bacterial growth inhibition assay. For example, an agar dilution assay identifies a substance that inhibits bacterial growth. Microtiter plates are prepared with serial dilutions of the test compound; adding to the preparation a given amount of growth substrate; and
10 providing a preparation of *Streptococcus* cells. Inhibition of growth is determined, for example, by observing changes in optical densities of the bacterial cultures.

- Inhibition of bacterial growth is demonstrated, for example, by comparing (in the presence and absence of a test compound) the rate of growth or the absolute growth of bacterial cells. Inhibition includes a reduction of one of the above
15 measurements by at least 20% (e.g., at least 25%, 30%, 40%, 50%, 75%, 80%, or 90%).

- Rodent (e.g., murine) and rabbit animal models of streptococcal infections are known to those of skill in the art, and such animal model systems are accepted for screening antibacterial agents as an indication of their therapeutic efficacy in
20 human patients. In a typical *in vivo* assay, an animal is infected with a pathogenic *Streptococcus* strain, e.g., by inhalation of *Streptococcus pneumoniae*, and conventional methods and criteria are used to diagnose the mammal as being afflicted with streptococcal pneumonia. The candidate antibacterial agent then is administered to the mammal at a dosage of 1-100 mg/kg of body weight, and the
25 mammal is monitored for signs of amelioration of disease. Alternatively, the test compound can be administered to the mammal prior to infecting the mammal with *Streptococcus*, and the ability of the treated mammal to resist infection is measured. Of course, the results obtained in the presence of the test compound should be compared with results in control animals, which are not treated with the test

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compound. Administration of candidate antibacterial agent to the mammal can be carried out as described below, for example.

Pharmaceutical Formulations

Treatment includes administering a pharmaceutically effective amount of a composition containing an antibacterial agent to a subject in need of such treatment, thereby inhibiting bacterial growth in the subject. Such a composition typically contains from about 0.1 to 90% by weight (such as 1 to 20% or 1 to 10%) of an antibacterial agent of the invention in a pharmaceutically acceptable carrier.

10 Solid formulations of the compositions for oral administration may contain suitable carriers or excipients, such as corn starch, gelatin, lactose, acacia, sucrose, microcrystalline cellulose, kaolin, mannitol, dicalcium phosphate, calcium carbonate, sodium chloride, or alginic acid. Disintegrators that can be used include, without limitation, micro-crystalline cellulose, corn starch, sodium starch
15 glycolate and alginic acid. Tablet binders that may be used include acacia, methylcellulose, sodium carboxymethylcellulose, polyvinylpyrrolidone (Povidone), hydroxypropyl methylcellulose, sucrose, starch, and ethylcellulose. Lubricants that may be used include magnesium stearates, stearic acid, silicone fluid, talc, waxes, oils, and colloidal silica.

20 Liquid formulations of the compositions for oral administration prepared in water or other aqueous vehicles may contain various suspending agents such as methylcellulose, alginates, tragacanth, pectin, kelgin, carrageenan, acacia, polyvinylpyrrolidone, and polyvinyl alcohol. The liquid formulations may also include solutions, emulsions, syrups and elixirs containing, together with the active
25 compound(s), wetting agents, sweeteners, and coloring and flavoring agents. Various liquid and powder formulations can be prepared by conventional methods for inhalation into the lungs of the mammal to be treated.

Injectable formulations of the compositions may contain various carriers such as vegetable oils, dimethylacetamide, dimethylformamide, ethyl lactate, ethyl

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carbonate, isopropyl myristate, ethanol, polyols (glycerol, propylene glycol, liquid polyethylene glycol, and the like). For intravenous injections, water soluble versions of the compounds may be administered by the drip method, whereby a pharmaceutical formulation containing the antibacterial agent and a physiologically acceptable excipient is infused. Physiologically acceptable excipients may include, for example, 5% dextrose, 0.9% saline, Ringer's solution or other suitable excipients. Intramuscular preparations, a sterile formulation of a suitable soluble salt form of the compounds can be dissolved and administered in a pharmaceutical excipient such as Water-for-Injection, 0.9% saline, or 5% glucose solution. A suitable insoluble form of the compound may be prepared and administered as a suspension in an aqueous base or a pharmaceutically acceptable oil base, such as an ester of a long chain fatty acid, (e.g., ethyl oleate).

A topical semi-solid ointment formulation typically contains a concentration of the active ingredient from about 1 to 20%, e.g., 5 to 10% in a carrier such as a pharmaceutical cream base. Various formulations for topical use include drops, tinctures, lotions, creams, solutions, and ointments containing the active ingredient and various supports and vehicles.

The optimal percentage of the antibacterial agent in each pharmaceutical formulation varies according to the formulation itself and the therapeutic effect desired in the specific pathologies and correlated therapeutic regimens. Appropriate dosages of the antibacterial agents can readily be determined by those of ordinary skill in the art of medicine by monitoring the mammal for signs of disease amelioration or inhibition, and increasing or decreasing the dosage and/or frequency of treatment as desired. The optimal amount of the antibacterial compound used for treatment of conditions caused by or contributed to by bacterial infection may depend upon the manner of administration, the age and the body weight of the subject and the condition of the subject to be treated. Generally, the antibacterial compound is administered at a dosage of 1 to 100 mg/kg of body weight, and typically at a dosage of 1 to 10 mg/kg of body weight.

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Example

Using the transposon-based mutagenesis methods described above, the *Streptococcus pneumonia* genome was mutagenized, and 23 genes were identified as being located within operons that are essential for survival of *Streptococcus pneumonia*. These genes are listed in Table 1, above, and their nucleic acid and amino acid sequences are represented by SEQ ID NOs:1-69, as shown in Figs. 1-23.

Now that each of these genes is known to be located within an operon that is essential for survival of *Streptococcus*, the polypeptides encoded by nucleic acids located within those operons can be used to identify antibacterial agents by using the assays described herein. Other art-known assays to detect interactions of test compounds with proteins, or to detect inhibition of bacterial growth also can be used with the nucleic acids located within operons containing the GEP genes, and gene products and homologs or orthologs thereof.

15 Other Embodiments

The invention also features fragments, variants, analogs, and derivatives of the GEP polypeptides described above that retain one or more of the biological activities of the GEP polypeptides, e.g., as determined in a complementation assay. Also included within the invention are naturally-occurring and non-naturally-occurring allelic variants. Compared with the naturally-occurring GEP gene, sequences depicted in Figs. 1-23, the nucleic acid sequence encoding allelic variants may have a substitution, deletion, or addition of one or more nucleotides. The preferred allelic variants are functionally equivalent to a GEP polypeptide, e.g., as determined in a complementation assay.

25 It is to be understood that, while the invention has been described in conjunction with the detailed description thereof, the foregoing description is intended to illustrate and not limit the scope of the invention, which is defined by the scope of the appended claims. Other aspects, advantages, and modifications are within the scope of the following claims.

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What is claimed is:

1. An isolated operon comprising a nucleotide sequence, or an allelic variant or homolog of the nucleotide sequence, encoding:
 - a gep103 polypeptide comprising the amino acid sequence of SEQ ID NO:1,
5 as depicted in Fig. 1;
a gep1119 polypeptide comprising the amino acid sequence of SEQ ID NO:4, as depicted in Fig. 2;
a gep1122 polypeptide comprising the amino acid sequence of SEQ ID NO:7, as depicted in Fig. 3;
10 a gep1315 polypeptide comprising the amino acid sequence of SEQ ID NO:10, as depicted in Fig. 4;
a gep1493 polypeptide comprising the amino acid sequence of SEQ ID NO:13, as depicted in Fig. 5;
a gep1507 polypeptide comprising the amino acid sequence of SEQ ID
15 NO:16, as depicted in Fig. 6;
a gep1511 polypeptide comprising the amino acid sequence of SEQ ID NO:19, as depicted in Fig. 7;
a gep1518 polypeptide comprising the amino acid sequence of SEQ ID NO:22, as depicted in Fig. 8;
20 a gep1546 polypeptide comprising the amino acid sequence of SEQ ID NO:25, as depicted in Fig. 9;
a gep1551 polypeptide comprising the amino acid sequence of SEQ ID NO:28, as depicted in Fig. 10;
a gep1561 polypeptide comprising the amino acid sequence of SEQ ID
25 NO:31, as depicted in Fig. 11;
a gep1580 polypeptide comprising the amino acid sequence of SEQ ID NO:34, as depicted in Fig. 12;
a gep1713 polypeptide comprising the amino acid sequence of SEQ ID NO:37 as depicted in Fig. 13;

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a gep222 polypeptide comprising the amino acid sequence of SEQ ID NO:40, as depicted in Fig. 14;

a gep2283 polypeptide comprising the amino acid sequence of SEQ ID NO:43, as depicted in Fig. 15;

5 a gep273 polypeptide comprising the amino acid sequence of SEQ ID NO:46, as depicted in Fig. 16;

a gep286 polypeptide comprising the amino acid sequence of SEQ ID NO:49, as depicted in Fig. 17;

a gep311 polypeptide comprising the amino acid sequence of SEQ ID
10 NO:52, as depicted in Fig. 18;

a gep3262 polypeptide comprising the amino acid sequence of SEQ ID NO:55, as depicted in Fig. 19;

a gep3387 polypeptide comprising the amino acid sequence of SEQ ID NO:58, as depicted in Fig. 20;

15 a gep47 polypeptide comprising the amino acid sequence of SEQ ID NO:61, as depicted in Fig. 21;

a gep61 polypeptide comprising the amino acid sequence of SEQ ID NO:64, as depicted in Fig. 22; or

a gep76 polypeptide comprising the amino acid sequence of SEQ ID NO:67,
20 as depicted in Fig. 23.

2. An isolated nucleic acid molecule comprising a nucleic acid sequence selected from the group consisting of:

(1) an operon comprising the sequence of SEQ ID NO:2, as depicted in Fig. 1, or degenerate variants thereof;

25 (2) an operon comprising the sequence of SEQ ID NO:2, or degenerate variants thereof, wherein T is replaced by U;

(3) nucleic acids complementary to (1) and (2);

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(4) fragments of (1), (2), and (3) that are at least 15 base pairs in length and which hybridize under stringent conditions to genomic DNA encoding the polypeptide of SEQ ID NO:1;

(5) an operon comprising the sequence of SEQ ID NO:5, as depicted in Fig. 2, or degenerate variants thereof;

(6) an operon comprising the sequence of SEQ ID NO:5, or degenerate variants thereof, wherein T is replaced by U;

(7) nucleic acids complementary to (5) and (6);

(8) fragments of (5), (6), and (7) that are at least 15 base pairs in length and which hybridize under stringent conditions to genomic DNA encoding the polypeptide of SEQ ID NO:4;

(9) an operon comprising the sequence of SEQ ID NO:8, as depicted in Fig. 3, or degenerate variants thereof;

(10) an operon comprising the sequence of SEQ ID NO:8, or degenerate variants thereof, wherein T is replaced by U;

(11) nucleic acids complementary to (9) and (10);

(12) fragments of (9), (10), and (11) that are at least 15 base pairs in length and which hybridize under stringent conditions to genomic DNA encoding the polypeptide of SEQ ID NO:7;

(13) an operon comprising the sequence of SEQ ID NO:11, as depicted in Fig. 4, or degenerate variants thereof;

(14) an operon comprising the sequence of SEQ ID NO:11, or degenerate variants thereof, wherein T is replaced by U;

(15) nucleic acids complementary to (13) and (14); and

(16) fragments of (13), (14), and (15) that are at least 15 base pairs in length and which hybridize under stringent conditions to genomic DNA encoding the polypeptide of SEQ ID NO:10;

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(17) an operon comprising the sequence of SEQ ID NO:14, as depicted in Fig. 5, or degenerate variants thereof;

(18) an operon comprising the sequence of SEQ ID NO:14, or degenerate variants thereof, wherein T is replaced by U;

5 (19) nucleic acids complementary to (17) and (18);

(20) fragments of (17), (18), and (19) that are at least 15 base pairs in length and which hybridize under stringent conditions to genomic DNA encoding the polypeptide of SEQ ID NO:13;

(21) an operon comprising the sequence of SEQ ID NO:17, as depicted in
10 Fig. 6, or degenerate variants thereof;

(22) an operon comprising the sequence of SEQ ID NO:17, or degenerate variants thereof, wherein T is replaced by U;

(23) nucleic acids complementary to (21) and (22);

(24) fragments of (21), (22), and (23) that are at least 15 base pairs in
15 length and which hybridize under stringent conditions to genomic DNA encoding the polypeptide of SEQ ID NO:16;

(25) an operon comprising the sequence of SEQ ID NO:20, as depicted in Fig. 7, or degenerate variants thereof;

(26) an operon comprising the sequence of SEQ ID NO:20, or degenerate
20 variants thereof, wherein T is replaced by U;

(27) nucleic acids complementary to (25) and (26);

(28) fragments of (25), (26), and (27) that are at least 15 base pairs in length and which hybridize under stringent conditions to genomic DNA encoding the polypeptide of SEQ ID NO:19;

25 (29) an operon comprising the sequence of SEQ ID NO:23, as depicted in Fig. 8, or degenerate variants thereof;

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(30) an operon comprising the sequence of SEQ ID NO:23, or degenerate variants thereof, wherein T is replaced by U;

(31) nucleic acids complementary to (29) and (30); and

(32) fragments of (39), (30), and (31) that are at least 15 base pairs in length and which hybridize under stringent conditions to genomic DNA encoding the polypeptide of SEQ ID NO:22;

(33) an operon comprising the sequence of SEQ ID NO:26, as depicted in Fig. 9, or degenerate variants thereof;

(34) an operon comprising the sequence of SEQ ID NO:26, or degenerate variants thereof, wherein T is replaced by U;

(35) nucleic acids complementary to (33) and (34);

(36) fragments of (33), (34), and (35) that are at least 15 base pairs in length and which hybridize under stringent conditions to genomic DNA encoding the polypeptide of SEQ ID NO:25;

(37) an operon comprising the sequence of SEQ ID NO:29, as depicted in Fig. 10, or degenerate variants thereof;

(38) an operon comprising the sequence of SEQ ID NO:29, or degenerate variants thereof, wherein T is replaced by U;

(39) nucleic acids complementary to (37) and (38);

(40) fragments of (37), (38), and (39) that are at least 15 base pairs in length and which hybridize under stringent conditions to genomic DNA encoding the polypeptide of SEQ ID NO:28;

(41) an operon comprising the sequence of SEQ ID NO:32, as depicted in Fig. 11, or degenerate variants thereof;

(42) an operon comprising the sequence of SEQ ID NO:32, or degenerate variants thereof, wherein T is replaced by U;

(43) nucleic acids complementary to (41) and (42);

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(44) fragments of (41), (42), and (43) that are at least 15 base pairs in length and which hybridize under stringent conditions to genomic DNA encoding the polypeptide of SEQ ID NO:31;

(45) an operon comprising the sequence of SEQ ID NO:35, as depicted in
5 Fig. 12, or degenerate variants thereof;

(46) an operon comprising the sequence of SEQ ID NO:35, or degenerate variants thereof, wherein T is replaced by U;

(47) nucleic acids complementary to (45) and (46); and

(48) fragments of (45), (46), and (47) that are at least 15 base pairs in
10 length and which hybridize under stringent conditions to genomic DNA encoding the polypeptide of SEQ ID NO:34;

(49) an operon comprising the sequence of SEQ ID NO:38, as depicted in Fig. 13, or degenerate variants thereof;

(50) an operon comprising the sequence of SEQ ID NO:38, or degenerate
15 variants thereof, wherein T is replaced by U;

(51) nucleic acids complementary to (49) and (50);

(52) fragments of (49), (50), and (51) that are at least 15 base pairs in length and which hybridize under stringent conditions to genomic DNA encoding the polypeptide of SEQ ID NO:37;

(53) an operon comprising the sequence of SEQ ID NO:41, as depicted in
20 Fig. 14, or degenerate variants thereof;

(54) an operon comprising the sequence of SEQ ID NO:41, or degenerate variants thereof, wherein T is replaced by U;

(55) nucleic acids complementary to (53) and (54);

(56) fragments of (53), (54), and (55) that are at least 15 base pairs in
25 length and which hybridize under stringent conditions to genomic DNA encoding the polypeptide of SEQ ID NO:40;

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(57) an operon comprising the sequence of SEQ ID NO:44, as depicted in Fig. 15, or degenerate variants thereof;

(58) an operon comprising the sequence of SEQ ID NO:44, or degenerate variants thereof, wherein T is replaced by U;

5 (59) nucleic acids complementary to (57) and (58);

(60) fragments of (57), (58), and (59) that are at least 15 base pairs in length and which hybridize under stringent conditions to genomic DNA encoding the polypeptide of SEQ ID NO:39;

(61) an operon comprising the sequence of SEQ ID NO:47, as depicted in
10 Fig. 16, or degenerate variants thereof;

(62) an operon comprising the sequence of SEQ ID NO:47, or degenerate variants thereof, wherein T is replaced by U;

(63) nucleic acids complementary to (61) and (62); and

(64) fragments of (61), (62), and (63) that are at least 15 base pairs in
15 length and which hybridize under stringent conditions to genomic DNA encoding the polypeptide of SEQ ID NO:46;

(65) an operon comprising the sequence of SEQ ID NO:50, as depicted in Fig. 17, or degenerate variants thereof;

(66) an operon comprising the sequence of SEQ ID NO:50, or degenerate
20 variants thereof, wherein T is replaced by U;

(67) nucleic acids complementary to (65) and (66);

(68) fragments of (65), (66), and (67) that are at least 15 base pairs in length and which hybridize under stringent conditions to genomic DNA encoding the polypeptide of SEQ ID NO:49;

25 (69) an operon comprising the sequence of SEQ ID NO:53, as depicted in Fig. 18, or degenerate variants thereof;

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(70) an operon comprising the sequence of SEQ ID NO:53, or degenerate variants thereof, wherein T is replaced by U;

(71) nucleic acids complementary to (69) and (70);

(72) fragments of (69), (70), and (71) that are at least 15 base pairs in length and which hybridize under stringent conditions to genomic DNA encoding the polypeptide of SEQ ID NO:52;

(73) an operon comprising the sequence of SEQ ID NO:56, as depicted in Fig. 19, or degenerate variants thereof;

(74) an operon comprising the sequence of SEQ ID NO:56, or degenerate variants thereof, wherein T is replaced by U;

(75) nucleic acids complementary to (73) and (74);

(76) fragments of (73), (74), and (75) that are at least 15 base pairs in length and which hybridize under stringent conditions to genomic DNA encoding the polypeptide of SEQ ID NO:55;

(77) an operon comprising the sequence of SEQ ID NO:59, as depicted in Fig. 20, or degenerate variants thereof;

(78) an operon comprising the sequence of SEQ ID NO:59, or degenerate variants thereof, wherein T is replaced by U;

(79) nucleic acids complementary to (77) and (78); and

(80) fragments of (77), (78), and (79) that are at least 15 base pairs in length and which hybridize under stringent conditions to genomic DNA encoding the polypeptide of SEQ ID NO:58;

(81) an operon comprising the sequence of SEQ ID NO:62, as depicted in Fig. 21, or degenerate variants thereof;

(82) an operon comprising the sequence of SEQ ID NO:62, or degenerate variants thereof, wherein T is replaced by U;

(83) nucleic acids complementary to (81) and (82);

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(84) fragments of (81), (82), and (83) that are at least 15 base pairs in length and which hybridize under stringent conditions to genomic DNA encoding the polypeptide of SEQ ID NO:61;

(85) an operon comprising the sequence of SEQ ID NO:65; as depicted in
5 Fig. 22, or degenerate variants thereof;

(86) an operon comprising the sequence of SEQ ID NO:65, or degenerate variants thereof, wherein T is replaced by U;

(87) nucleic acids complementary to (85) and (86);

(88) fragments of (85), (86), and (87) that are at least 15 base pairs in
10 length and which hybridize under stringent conditions to genomic DNA encoding the polypeptide of SEQ ID NO:66;

(89) an operon comprising the sequence of SEQ ID NO:68, as depicted in Fig. 23, or degenerate variants thereof;

(90) an operon comprising the sequence of SEQ ID NO:68, or degenerate
15 variants thereof, wherein T is replaced by U;

(91) nucleic acids complementary to (89) and (90); and

(92) fragments of (89), (90), and (91) that are at least 15 base pairs in length and which hybridize under stringent conditions to genomic DNA encoding the polypeptide of SEQ ID NO:67.

20 3. An isolated operon from *Streptococcus* comprising a nucleotide sequence that is at least 85% identical to a nucleotide sequence selected from the group consisting of

SEQ ID NO:2;

SEQ ID NO:5;

25 SEQ ID NO:8;

SEQ ID NO:11;

SEQ ID NO:14;

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5 SEQ ID NO:17;
SEQ ID NO:20;
SEQ ID NO:23;
SEQ ID NO:26;
SEQ ID NO:29;
SEQ ID NO:32;
SEQ ID NO:35;
SEQ ID NO:38;
SEQ ID NO:41;
10 SEQ ID NO:44;
SEQ ID NO:47;
SEQ ID NO:50;
SEQ ID NO:53;
SEQ ID NO:56;
15 SEQ ID NO:59;
SEQ ID NO:62;
SEQ ID NO:65; and
SEQ ID NO:68.

4. An isolated nucleic acid molecule that is at least 15 base pairs in length
20 and hybridizes under stringent conditions to a nucleotide sequence selected from
the group consisting of

SEQ ID NO:2;
SEQ ID NO:5;
SEQ ID NO:8;
25 SEQ ID NO:11;
SEQ ID NO:14;
SEQ ID NO:17;
SEQ ID NO:20;
SEQ ID NO:23;

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SEQ ID NO:26;
SEQ ID NO:29;
SEQ ID NO:32;
SEQ ID NO:35;
5 SEQ ID NO:38;
SEQ ID NO:41;
SEQ ID NO:44;
SEQ ID NO:47;
SEQ ID NO:50;
10 SEQ ID NO:53;
SEQ ID NO:56;
SEQ ID NO:59;
SEQ ID NO:62;
SEQ ID NO:65; and
15 SEQ ID NO:68.

5. A vector comprising an operon of claim 1.

6. A vector comprising a nucleic acid molecule of claim 2.

7. An expression vector comprising an operon of claim 1 operably linked to a nucleotide sequence regulatory element that controls expression of said operon.

20 8. An expression vector comprising a nucleic acid molecule of claim 2, wherein said nucleic acid molecule is operably linked to a nucleotide sequence regulatory element that controls expression of said nucleic acid.

9. A host cell comprising an exogenously introduced operon of claim 1.

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10. A host cell comprising an exogenously introduced nucleic acid molecule of claim 2.

11. A host cell of claim 9, wherein the cell is a yeast or bacterium.

12. A host cell of claim 10, wherein the cell is a yeast or bacterium.

5 13. A genetically engineered host cell comprising an operon of claim 1 operably linked to a heterologous nucleotide sequence regulatory element that controls expression of the operon in the host cell.

14. A host cell of claim 13, wherein the cell is a yeast or bacterium.

10 15. A genetically engineered host cell comprising a nucleic acid molecule of claim 2 operably linked to a nucleotide sequence regulatory element that controls expression of the nucleic acid in the host cell.

16. A host cell of claim 15, wherein the cell is a yeast or bacterium.

17. An isolated operon comprising a nucleotide sequence encoding a polypeptide comprising an amino acid sequence selected from the group consisting
15 of:

the amino acid sequence of SEQ ID NO:1, as depicted in Fig. 1;
the amino acid sequence of SEQ ID NO:4, as depicted in Fig. 2;
the amino acid sequence of SEQ ID NO:7, as depicted in Fig. 3;
the amino acid sequence of SEQ ID NO:10, as depicted in Fig. 4;
20 the amino acid sequence of SEQ ID NO:13, as depicted in Fig. 5;
the amino acid sequence of SEQ ID NO:16, as depicted in Fig. 6;
the amino acid sequence of SEQ ID NO:19, as depicted in Fig. 7;
the amino acid sequence of SEQ ID NO:22, as depicted in Fig. 8;

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the amino acid sequence of SEQ ID NO:25, as depicted in Fig. 9;
the amino acid sequence of SEQ ID NO:28, as depicted in Fig. 10;
the amino acid sequence of SEQ ID NO:31, as depicted in Fig. 11;
the amino acid sequence of SEQ ID NO:34, as depicted in Fig. 12;
5 the amino acid sequence of SEQ ID NO:37, as depicted in Fig. 13;
the amino acid sequence of SEQ ID NO:40, as depicted in Fig. 14;
the amino acid sequence of SEQ ID NO:43, as depicted in Fig. 15;
the amino acid sequence of SEQ ID NO:46, as depicted in Fig. 16;
the amino acid sequence of SEQ ID NO:49, as depicted in Fig. 17;
10 the amino acid sequence of SEQ ID NO:52, as depicted in Fig. 18;
the amino acid sequence of SEQ ID NO:55, as depicted in Fig. 19;
the amino acid sequence of SEQ ID NO:58, as depicted in Fig. 20;
the amino acid sequence of SEQ ID NO:61, as depicted in Fig. 21;
the amino acid sequence of SEQ ID NO:64, as depicted in Fig. 22; and
15 the amino acid sequence of SEQ ID NO:67, as depicted in Fig. 23.

18. An isolated polypeptide encoded by a nucleic acid located within an operon comprising a nucleic acid sequence selected from the group consisting of SEQ ID NO: 2, 5, 8, 11, 14, 17, 20, 23, 26, 29, 32, 35, 38, 41, 44, 47, 50, 53, 56, 59, 62, 65, and 68.

20 19. An isolated polypeptide, said polypeptide being encoded by an operon of claim 1.

20. An isolated polypeptide, said polypeptide being encoded by a nucleic acid molecule of claim 2.

21. An isolated polypeptide, said polypeptide being encoded by an
25 operon of claim 3.

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22. A method for identifying an antibacterial agent, the method comprising:

(a) contacting a test compound with a polypeptide, or a homolog of a polypeptide, encoded by a nucleic acid sequence located within an operon comprising a GEP gene selected from the group consisting of gep103, gep1119, gep1122, gep1315, gep1493, gep1507, gep1511, gep1518, gep1546, gep1551, gep1561, gep1580, gep1713, gep222, gep2283, gep273, gep286, gep311, gep3262, gep3387, gep47, gep61, and gep76; and

(b) detecting binding of the test compound to the polypeptide, wherein binding indicates that the test compound is an antibacterial agent.

23. The method of claim 22, further comprising:

(c) determining whether a test compound that binds to the polypeptide inhibits growth of bacteria, relative to growth of bacteria cultured in the absence of a test compound that binds to the polypeptide, wherein inhibition of growth indicates that the test compound is an antibacterial agent.

24. The method of claim 22, wherein the polypeptide is selected from the group consisting of gep103, gep1119, gep1122, gep1315, gep1493, gep1507, gep1511, gep1518, gep1546, gep1551, gep1561, gep1580, gep1713, gep222, gep2283, gep273, gep286, gep311, gep3262, gep3387, gep47, gep61, and gep76.

25. The method of claim 22, wherein the test compound is immobilized on a substrate, and binding of the test compound to the polypeptide is detected as immobilization of the polypeptide on the immobilized test compound.

26. The method of claim 25, wherein immobilization of the polypeptide on the test compound is detected in an immunoassay with an antibody that specifically binds to the polypeptide.

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27. The method of claim 22, wherein the test compound is selected from the group consisting of polypeptides and small molecules.

28. The method of claim 22, wherein:

(a) the polypeptide is provided as a fusion protein comprising the
5 polypeptide fused to (i) a transcription activation domain of a transcription factor or
(ii) a DNA-binding domain of a transcription factor; and

(b) the test compound is a polypeptide that is provided as a fusion protein
comprising the test polypeptide fused to (i) a transcription activation domain of a
transcription factor or (ii) a DNA-binding domain of a transcription factor, to
10 interact with the fusion protein; and

(c) binding of the test compound to the polypeptide is detected as
reconstitution of a transcription factor.

29. An antibody that specifically binds to a GEP polypeptide of claim 19.

30. An antibody of claim 29, wherein the antibody is a monoclonal
15 antibody.

31. A method for identifying an antibacterial agent, the method comprising:

(a) contacting a polypeptide encoded by a nucleic acid located within an
operon comprising a GEP gene with a test compound;

(b) detecting a decrease in function of the polypeptide contacted with the
20 test compound; and

(c) determining whether a test compound that decreases function of a
contacted polypeptide inhibits growth of bacteria, relative to growth of bacteria
cultured in the absence of a test compound that decreases function of a contacted
polypeptide, wherein inhibition of growth indicates that the test compound is an
25 antibacterial agent.

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32. The method of claim 31, wherein the polypeptide is selected from the group consisting of gep103, gep1119, gep1122, gep1315, gep1493, gep1507, gep1511, gep1518, gep1546, gep1551, gep1561, gep1580, gep1713, gep222, gep2283, gep273, gep286, gep311, gep3262, gep3387, gep47, gep61, and gep76.

5 33. The method of claim 31, wherein the test compound is selected from the group consisting of polypeptides and small molecules.

34. A method for identifying an antibacterial agent, the method comprising:

(a) contacting a nucleic acid comprising an operon containing a gene encoding a GEP polypeptide with a test compound, wherein the GEP polypeptide is
10 selected from the group consisting of gep103, gep1119, gep1122, gep1315, gep1493, gep1507, gep1511, gep1518, gep1546, gep1551, gep1561, gep1580, gep1713, gep222, gep2283, gep273, gep286, gep311, gep3262, gep3387, gep47, gep61, and gep76; and

(b) detecting binding of the test compound to the nucleic acid, wherein
15 binding indicates that the test compound is an antibacterial agent.

35. The method of claim 34, further comprising:

(c) determining whether a test compound that binds to the nucleic acid inhibits growth of bacteria, relative to growth of bacteria cultured in the absence of the test compound that binds to the nucleic acid, wherein inhibition of growth
20 indicates that the test compound is an antibacterial agent.

36. The method of claim 34, wherein the test compound is selected from the group consisting of polypeptides and small molecules.

37. An isolated nucleic acid or an allelic variant thereof encoding:
a gep1493 polypeptide comprising the amino acid sequence of SEQ ID

25 NO:13, as depicted in Fig. 5;

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a gep1507 polypeptide comprising the amino acid sequence of SEQ ID NO:16, as depicted in Fig. 6;

a gep1546 polypeptide comprising the amino acid sequence of SEQ ID NO:25, as depicted in Fig. 9;

5 a gep273 polypeptide comprising the amino acid sequence of SEQ ID NO:46, as depicted in Fig. 16;

a gep286 polypeptide comprising the amino acid sequence of SEQ ID NO:49, as depicted in Fig. 17; or

10 a gep76 polypeptide comprising the amino acid sequence of SEQ ID NO:67, as depicted in Fig. 23.

38. An isolated nucleic acid comprising a sequence selected from the group consisting of:

(1) SEQ ID NO:14, as depicted in Fig. 5, or degenerate variants thereof;

15 U;
(2) SEQ ID NO:14, or degenerate variants thereof, wherein T is replaced by

(3) nucleic acids complementary to (1) and (2);

(4) fragments of (1), (2), and (3) that are at least 15 base pairs in length and which hybridize under stringent conditions to genomic DNA encoding the polypeptide of SEQ ID NO:13;

20 (5) SEQ ID NO:17, as depicted in Fig. 6, or degenerate variants thereof;

(6) SEQ ID NO:17, or degenerate variants thereof, wherein T is replaced by U;

(7) nucleic acids complementary to (5) and (6);

25 (8) fragments of (5), (6), and (7) that are at least 15 base pairs in length and which hybridize under stringent conditions to genomic DNA encoding the polypeptide of SEQ ID NO:16;

(9) SEQ ID NO:26, as depicted in Fig. 9, or degenerate variants thereof;

(10) SEQ ID NO:26, or degenerate variants thereof, wherein T is replaced by U;

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- (11) nucleic acids complementary to (9) and (10);
- (12) fragments of (9), (10), and (11) that are at least 15 base pairs in length and which hybridize under stringent conditions to genomic DNA encoding the polypeptide of SEQ ID NO:25;
- 5 (13) SEQ ID NO:47, as depicted in Fig. 16, or degenerate variants thereof;
- (14) SEQ ID NO:47, or degenerate variants thereof, wherein T is replaced by U;
- (15) nucleic acids complementary to (13) and (14);
- (16) fragments of (13), (14), and (15) that are at least 15 base pairs in
- 10 length and which hybridize under stringent conditions to genomic DNA encoding the polypeptide of SEQ ID NO:46;
- (17) SEQ ID NO:50, as depicted in Fig. 17, or degenerate variants thereof;
- (18) SEQ ID NO:50, or degenerate variants thereof, wherein T is replaced by U;
- 15 (19) nucleic acids complementary to (i) and (j);
- (20) fragments of (i), (j), and (k) that are at least 15 base pairs in length and which hybridize under stringent conditions to genomic DNA encoding the polypeptide of SEQ ID NO:49;
- (21) SEQ ID NO:68, as depicted in Fig. 23, or degenerate variants thereof;
- 20 (22) SEQ ID NO:68, or degenerate variants thereof, wherein T is replaced by U;
- (23) nucleic acids complementary to (21) and (22); and
- (24) fragments of (21), (22), and (23) that are at least 15 base pairs in length and which hybridize under stringent conditions to genomic DNA encoding
- 25 the polypeptide of SEQ ID NO:67.

39. A method for identifying an antibacterial agent, the method comprising:

- (a) contacting a test compound with a polypeptide, or a homolog of a polypeptide, encoded by a nucleic acid sequence located within an operon comprising a B-yneS gene; and

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(b) detecting binding of the test compound to the polypeptide, wherein binding indicates that the test compound is an antibacterial agent.

40. The method of claim 39, further comprising:

- (c) determining whether a test compound that binds to the polypeptide
- 5 inhibits growth of bacteria, relative to growth of bacteria cultured in the absence of a test compound that binds to the polypeptide, wherein inhibition of growth indicates that the test compound is an antibacterial agent.

sep103

Fig. 1

(SEQ ID NO: 2) 1 TCGTGATTTTGGAGAAAGTTTATTACAGATAAAAGAGTCTAAGGAAAAAATCCATTTGATATTTTCTTCTATAAAATAGATAAAATGGTACAATA 100
ACGACTAAAAACCTCTTTCAAATAATCTCTATTTTTCAGATTCTTTTTTAAGTAAACTATAAAAGAAGATATTTATCTATTTTACCATGTTAT

(SEQ ID NO: 3)

101 ATAAATGAGGTAAATAGGATCAGATTAGATAAATATTTAAAGTATCCCAATTATCAAGCGTCCTACAGTCCCAAGGAAGTAGCAGATAAAGGTAGA 300
TATTTAACTCCATTATCTCTACTCTAATCTATTTATAAAATTTTCATAGCGCTTAATAGTTCGCAGCATGTCAGCGTTTCTTTCATCGTCTATTTCCATCT

(SEQ ID NO: 1) 1 M R L D K Y L K V S R I I K R R T V A K E V A D K G R 27

201 ATCAAGGTTAATGGAATCTTGGCCAAAAGTTCAACGGACTTGAAAGTTAATGACCAAGTTGAAATTCGCTTGGCAATAAGTTGCTGCTTGTAAAAGTAC 300
TAGTTCCAATTACCTTAGAACCGGTTTTCAAGTTGCCCTGAACTTTCAATTACTGGTTCAACTTAAGCGAAACCGTTATTCACGACGAACATTTTCATG

28 I K V N G I L A K S S T D L X V N D Q V E I R F G N K L L L V K V L 61

301 TAGAGATGAAAGATAGTACAAAAAAGAAGATGCAGCAGGAATGTATGAAATTTATCAGTGAACACCGGTAGAAGAAAAATGTCTAAAAATATTGTACAAT 400
ATCTCTACTTTCTATCATGTTTTTTCTTCTACGTGTCCTTACATACCTTTAATAGTCACTTTGTGCCCATCTTCTTTTACAGATTTTATAACATGTTA

62 E M K D S T K X E D A A G M Y E I I S E T R V E E N V . 89

gsep1313

Fig. 2

(SEQ ID NO: 5) 1 GAAATCCGTTTCAAATGTGACTGTAGCCATGAACGCTTTATGAACGCTCTTCCAGCTTCCAAGCTCAGACTTACAGGAAATGAAGAGGGAAGACCAAG 100
(SEQ ID NO: 6) CTTTAGGCAAAAGTTACACTGACATCGGTACTTGGAAATATCTTGGAGAACGGTTCGGAAGGTTGAGTCTGAAATGTCCTTACTTTCTCTTCTGGTGC

101 GGGCAGAAATCACTTGTCAATTCGCCAACTACTTACAACCTTGTATGAAAGGACCTGGAGGAACTCATTCGTGACAAATCTTAAATACACCTTTTATCA 200
CCCGTCTTATGTGAACAGTTAAGACGGTTTGTATGAATGTTGAAACTACTTTTCTGGACCTCTTGAGTAAGCACTGTTTAGAATTATGTGGAAATACT

(SEQ ID NO: 4) 1 M K R T W R N S F V T N L N T P F M I 19

201 TTGGCAATATTGAGATTCCCAATCGTACCGTTTATAGCGCTATGGCTGGCGTACCAACTCAGCCTTTCGTACCATCGCAAAAGAGCTCGGAGCTGGACT 300
AACCGTTATAACTCTAAGGGTTAGCATCGCAAAATCGCGATACCGACCGCACTGGTTGAGTCGGAAGCATGGTAGCGTTTCTCGAGCCTCGACCTGA

20 G N I E I P N R T V L A P M A G V T N S A P R T I A K E L G A G L 52

301 CGTTGTAAATGGAAATGGTCTCTGACAAGGGAATCCAATACAACAAGAAACCCCTGCATATGCTTCATATCGATGAGGGGGAACCCCTGTCTCTATC 400
GCAACATTACCTTTACAGAGACTGTTCCCTTAGGTTATGTTGCTTTTGGGAGGTATACGAATATAGCTACTCCCGCTTTTGGGACAGAGATAG

53 V V M E M V S D K G I Q Y N N E K T L H M L N I D E G E N P V S I 85

401 CAACTTTTGGTAGCGATGAAGACAGCCTAGCAGCGCAGCAGAAATTCATCCAAGAAAACCAAGACCGATATCGTGGATATCAACATGGGCTGCCCTG 500
GTTGAAAACCATCGCTACTTCTGTGGATCGTGGCGTCTCTAAGTAGGTTCTTTTGTGTTCTGGCTATAGCAGCTATAGTTGTACCCGACGGGAC

86 O L F G S D E D S L A R A A E F I O E N T K T D I V D I N H C C P V 119

0

501 TCAACAAATCGTGAAGAAGCAAGCTGGAGCTATGTGGCTCAAGGATCCTGACAGATCTACTCTATCATCAACAGGTCCAGTCTGTCTTATATCCC 600
AGTTGTTTATGCACTTCTTCTCTGACCTCGATACCCGAGTCTTAGGACTGTTCTAGATGAGATAGTAGTTGTTCCAGGTGAGACAGCACTATAGGG

120 N K I V K N E A G A M M L K D P D K I Y S I I N K V Q S V L D I P 152

601 ACTTACTGTCAAAATGGTACCGGCTGGGCGGACCACTCTGCGCAGTAGAAAATGCGCTCGCTGCTGAGCGTCCAGGTGTTTCTGCCCTCGCCATGCAT 700
TGAATGACAGTTCACGATGGCGGACCGGCTGGTAGAGACCGTCACTCTTACGGGAGCGAGCACTCGACGTCCACAAAGACGGGAGCGGTACGTA

153 L T V K M R T G W A D P S L A V E N A L A A E A A G V S A L A M H 185

701 GGGCGTACCGGTGAACAAATGTATACTGGCCACGAGACCTTGAGACCTTTACAAGGTTGCCAAGCTCTAACCAAGATTCCATTATCGCCAAAGGTC 800
CCGGCATGGGCACTTGTATCATATGACCGGTGGCTCTGGAACCTTGGGAAATGTTCCAAGGGTTCGAGATTGGTCTAAGGTAAGTAGCGGTGCCAC

186 G R T R E Q M Y T G H A D L E T L Y K V A Q A L T K I P F I A N G D 219

801 ATATCCGTACTGTCCAAGAGCCAAGCAACGATCGAAGAAGTGGTGTGACGAGTCATGATTGGCCGAGCTGCCATGGGAAATCCTTACCTCTTCAA 900
TATAGGCATGACAGGTTCTCGGTTCTTGGTAGCTTCTCAACCAGCACTGCGTCAGTACTAACCGGCTCGACGGTACCTTTAGGAATGGAGAAGTT

220 I R T V Q E A K O R I E E V G A D A V M I G R A A M G N P Y L F N 252

901 CCAATCAACCATTACTTTGAACACGAGAAATCTACCTGATTTGACCTTTGAAGACAGATCAAGATCCCTACGAACACTTGAAGCGATTGATTAACT 1000
CGTTAGTTGGTAATGAACCTTTCTCTCTTAGGATGAGCTAACTGGAACTCTGTTCTACTTCTAGCGGATGCTTGTGAACCTTCTTAACCTAATTG

253 O I N H Y F E T G E I L P D L T F E D K M K I A Y E H L K R L I N 285

1001 CTCAAAGGAGAAAAGCTCGCAGTTCTGAAATTCGCGGCTCGCTCTCACTATCTCGTGGAAACATCTGGCGCTGCCAAACTCCGTGGAGCCATTTCCG 1100
GAGTTCTCTTTTGCAGCTCAAGCACTAAGCGCGCGAGCGAGGAGTGATAGAGGCACCTTGTAGACCGGACGGTTTGAAGCACTCGGTAAAGCG

286 L K G E N V A V R E F R G L A P H Y L R G T S G A A K L R G A I S O 319

1101 AAGCTAGACCCCTAGCAGAGATTGAAGCCCTCTTGAATTCGAGAAGGCTTAATAGTTTAAACCCGTAACCTCTCTTAAGAGTCTCTTGAATGCCGCCA 1200
TTCCATCGTGGATCGTCTAAGTTCCGGAGAACGTTAACTCTTCCGAATTATCAAAATTTGGGCATTGAGAGAAATTTCTCAGAGAACTTACGGCGGT

320 A S T L A E I E A L L O L E K A 336

gopl122

Fig. 3 (Sheet 1 of 2)

(SEQ ID NO: 8) 1 AAGGACGAGCTGGAGTTTCCCTCATATTTTCAATAGTTTATTAGCTACAGTTGAGCACTTCAGAAAAATCAAATTTCTTCAAGTTCTCTCTTA 100
(SEQ ID NO: 9) TCCGTGCTCGACCTTCAAAAGGAGTATAAAAAAGTTATCAATAATCGATGTCGAACCTGTTGAAGTCTTTTAGTTTAAAGAAAGTTCAAGAGAAGAT

101 TAGTAGATTTTGAATCCCTTTTGAAGTAGTTTCTGAGTCAGCACATAAGGACCTTGTCTCCTGAAAGTTGATTGGTATTCATAGCATAGCATAAAGCTTA 200
ATCATCTAAAACCTTAGGGAAAACTCGATCAAGACTCAGTCGTGTATTCTCGGAAACAGAGGACTTTCAACTAACCATAACTACTATCGTATTCCGAT

201 CTGACCATCATTAAATCCACTTATCTTTTAAAGATTAGCAATACTTGAGAAACGATGTTTTATCAATATCGTATTTTTCAGATATTTCTGACTTTCT 300
GACTGGTAGTAATTAGGTGAATAGAGAAATTTCTAATCGTTATTCGAACCTTTCTGTACAAAAATAGTTATAGCATAAAAAGTCTATAAGAGAGTGAAGA

301 TTTTCAGTGGTGTCTTAAAGGATAAGTGGTAGAGGGCCAGATTCTTACCATAGAAAAATTCAGCAAGTCTTGAATCTCTTTCAATCTCTCTCGCTTA 400
AAAAGTCACGCACGAAATTTCTATTACCATCTCCCGGTCTAAGAAATGGTATTCTTTAACTCGTTTCAGAACTTAGAGAAAGTTAAGGAGAGCGAAT

401 TCACCTTATCTCTCGATAACATAAAACGAACAAATGTATCTCGGTGATATAGCAATTTGTCCGCAATATCAAGCTCCATCAGATAGAGTCTTTTCTT 500
AGTGGAAATAGAGAGCTATTGTATTTCTTGTAAATAGAGCCACTATATCGTAAACAGCGGTAAATAGTTTCAGAGTAGCTCTATCTCAGAAAAAAGAA

501 TTCAAGTTTTGTGATTTTCTAGCTCTATTATAACTCAAAATGTGATAAGATAGGGGTATGAATCTGAAAGTGAAAAAATACCATTAATAATCAAG 600
AAGTTCAAAACACTAAAGTATCGAGATAATATTGAGTTTACACTATTTCTATCCCATACTTAGACTTTCACTTTGTTTTTATGGTAAATTTTAGTTT

(SEQ ID NO: 7) 1 M N L K V K Q K I P L K I K 14

601 CGCATGGGAATTAACGGTGAGGGAATCGGCTTTTACAAAAACATTAGTCTTTGTACAGGAGCTCTCAAGGCGAAGATATCTATTGTGAGATTACTT 700
CGGTACCTTAATTCGCACTCCCTTAGCCGAAATGGTTTTGTAAATCAGAAACATGGTCTCGAGAGTTTCCGCTCTATAGATAACAGTCTAATGAA

15 R M G I N G E G I C F Y Q K T L V F V P G A L K G E D I Y C Q I T S 48

701 CTATTAGACGCAACTTTGTTGAAGCAAAATTAAGTCAAGGTCACCAAGAGTCTAAATTTGAAATTTGTCATCTTGTACTATTATAATGAATGCCGAGG 800
GATAATCTGGCTTGAAACAACTTCGTTTAAATGACTTCCAGTGTCTTTCAGATTAAAGCTTAACACGGTAGAACAATGATAAATATTACTTACGCTCC

49 I R R N F V E A K L L K V N K K S K F R I V P S C T I Y N E C G G 81

801 CTGCCAAATCATGCACCTGCATTATGATAAGCAGCTCGAGTTCAAGACCGCACTTACTTCATCAAGCGCTGAAAAAATTTGCTCTGCAGGATATGAAAT 900
GACGGTTTAGTACGTGAGCTAATACTATTCTGACCTCAAGTTCTGCTGAATGAAGTAGTTCCGCACTTTTAAACGAGGAGCTCTATACTTTTA

82 C Q I M H L H Y D K O L E F K T D L L H O A L K K F A P A G Y E N 114

901 TATGAAATTCCTCCAACTATTGGAATGCAGGAACCAAAATATTACAGAGCTAAGTTACAAATTCAGACTCGAAAAATTTAAAAATCAGGTCAAGCGCGGCT 1000
ATACTTTAAGCAGGTTGATAACCTTACGTCTTGGTTTTATAATGCTCGATTCAATGTTAAAGTCTGAGCTTTTAAATTTTAGTCCAGTTCCGCCCCGA

115 Y E I R P T I G M Q E P K Y Y R A K L O F O T R K F K N O V K A G L 148

1001 TATATGCACAAAATCTCACTATTAGTAGAGTTGAAAGACTGCTGGTACAAGATAAGGAAACCAAGTGATTGCTAATCGCTTAGCAGAAATTAATTAC 1100
ATATACGTGTTTGAAGTATGATAATCATCTCAACTTTCTGACGGACCATGTTCTATTCTTTGGGTTCACTAACGATTAGCGAATCGTCTTAATGAATG

149 Y A Q N S H Y L V E L K D C L V Q D K E T Q V I A N R L A E L L T 181

1101 TTATCACCAGATTCCAATCACGATGAGAGAAAAAGTTTAGGTGTCTGACTATTATGGTCCGACGCGGAGAAAGACCGGACAGGTTCAAGATTATTAT 1200
AATAGTGGTCTAAGGTTAGTGCCTACTCTCTTTCAAGATCCACAGGCATGATAATACCAGGCTGCGGCTCTTTCTGGCTGTCCAGCTTAATAATAA

182 Y H Q I P I T D E R K V L G V R T I M V R R A R K T G O V Q I I I 214

1201 GTTACAAACCGCCAGCTTAATTTAACTCAATTCGTAAGAGGTTGGTTAAAGATTCCAGAAAGTTGTGACAGTAGCTGTTAATACAAATACAGCTAAAA 1300
CAATGTTTGGCGGTGAAATTAATTTGAGTTAAACATTTCTCAACCAATTTCTAAAGGCTCTTCAACACTGTATCGACAAATTTATGTGCAATTTT

215 V T N R O L N L T O L V K E L V K D F P E V V T V A V N T N T A K T 248

1301 CCAGTGCATATATGGTGAAGAGACAGAGATTATCGGGGCAAGAGATATTCAAGAAAGTGTACTCAATTATGAATTTTCACTATCCGCTCGAGCTTT 1400
GGTCACTCTATATACCACTTTCTGTCTTAATAGACCCGCTCTCTCAATAGTTCTTCCACATGAGTTAACTTAAAGTGATAGGGAGCTCGAAA

Fig. 3 (Sheet 2 of 2)

249 S E I Y G E K T E I I M G O E S I O E G V L N Y E F S L S P R A F 281

1401 TTATCAACTAAATCCTGAGCAAAACAGAACTCCTCTATAGCGAAGCAGTAAAGCGCTGGATGTTGATAAAGAAGACCATTTCATTGACGCTTATTGTGGA 1500
AATAGTTGATTTAGGACTCGTTTGTCTTCAGGAGATATCGCTTCTCATTTCGCGACCTACAACTATTCTCTGGTAAACTAAGTGGAAATAACACT

282 Y Q L N P E O T E V L Y S E A V R A L D V D K E D H L I D A Y C G 314

1501 GTTGGACGATTGGATTTCCTTTGCAAGAAAGTAAAAACACTCAGAGGTATGGATATTATTCAGAAGCTATTGAAGATGCCAAGCGAAATGCTAAAA 1600
CAACTTGTCTAACTAAACGGAAACGTTTCCTTCACTTTTGTGAGTCTCCATACCTATAATAAGGTCTTCGATAACTTCTACGGTTTCGTTTACGATTTT

315 V G T I G F A F A K K V K T L R G M D I I F E A I E D A K R N A K R 348

1601 GAATGGGATTGACAATACTCATTATGAAGCTGGAAACGAGAGAGATTATTCCTCTTGGTACAAGCAAGCTACCGAGCAGATGCTTTGATTGTTGA 1700
CTTACCTTAACTGTTATGAGTAATCTTCGACCTTGGCGTCTTCTCTAATAAGGAGCAACCATGTTCTTCGATGGCTCGTCTACGAACTAAACACT

349 M C F D N T H Y E A G T A E E I I P R W Y K E G Y R A D A L I V D 381

1701 CCCACCAGGTACAGGTCTGGATGATAAGTTATTAGATACTATTCTACTATGTACCAGAAAAAATGGTTTATATTCTTGTATGTTTCGACCTTGGCT 1800
GGTGGTGCATGTCCAGACCTACTATTCAATAATCTATGATAAAGATGAATACATGCTCTTTTACCAATATAAAGAACATTACAAGCTGGAAACGA

382 P P R T G L D D K L L D T I L T Y V P E K M V Y I S C N V S T L A 414

1801 CGTGATTGGTACGCTTAGTAGAAGTCTATGATCTTCATTATATCCAGTGGTGGATATGTTCCACATACAGCTCGAAGTGAAGCTGTTGTAATAATTAA 1900
GCACTAAACCATGCCAATCATCTTCAGATACTAGAAGTAATATAGGTGAGCCAGCTATACAAGGGTGTATGTGAGCTTGACTTCGACAACTTTTAATT

415 R D L V R L V E V Y D L H Y I O S V D M F P H T A R T E A V V K L I 448

1901 TAACAAAAGTTTAAAAAGTAGTTGACAAAGTTTCAAAAGACTGTATAATAGTAAGAGTTGAAAAAACAACCTCAGGTTCGTTGGTCAAGGGGTAAAGAC 2000
ATTGTTTTCAAAATTTTTCATCAACTGTTTCAAACTTTTCTGACATATTATCTTCTCAACTTTTATTGTTGAGTCCAGCAACCACTTCCCAATTCTG

449 T K V • 452

2001 ACCGCTTTTCACGGCGGTAAACACGGGTTGGAATCCCGTACGGACTATGGTATGTTGCGGTTGGAACTTGTATGAAAACTTTA 2084
TGCGGAAAAGTGCGCCATTGTGCCCCAAGCTTAGGGCATGCTGATACCATACAACGCCAACCTTGTGAACTACTTTTTGAAAT

g9p1315

Fig. 4

(SEQ ID NO:11) ¹ AAGAGCTCCTTTCTTTTATTTATCTTAGCAAAATTCCTCMAATTAGTAGTAGCATAGCCTGTTTGTACTGGCTAAAAACAGGCTATTCAAATTCAG 100
(SEQ ID NO:12) TTCTCGAGGAAGAAAAATAAATAGAATCGTTAAAGGGAGTTTAAATCGATCATCGTATCGGACAAACATGACCGATTTTTGTCCGATAAAGTTTAAGTC

101 TTTCAGACCATCTAGCATCGAAAAATCTGTTATAATAATGGAAAAGGAGAGCGCATGCCAAGATTTTATAAGAGATGATCAGGTCATTCTGTCAA 200
AAAGTCTGGTAGATCGTACCTTTTAGACAATATTATTACCTTTCTCTTCGCTACGTTTCTAAAAATAATTATCTTCTACTAGTCCAGTAAGCAGTT

(SEQ ID NO:10) ¹ M H K I L L I E D D O V I R Q 15

201 CAGATTGGGAAATGCTCTCTCAATCGGATTTNAAGTGGTCTGGTAGAAGACTTATGGAAGTTTGAAGTCTATTGTTCAAGTCGGAACTTCATCTGG 300
GTCTAACCTTTTACGAGAGACTTACCTTAAATTCACGAGGACCATCTTCTGAAATACCTTCAAAACTCAGATAAACAGTACGCTTCGAGTAGACC

16 Q I G K M L S E W G F X V V L V E D P M E V L S L F V Q S E P H L V 49

301 TCCTCATGATATTGGTTTGCCTTGTTTAATGGTTATCACTGGTTCAGGAAATCCGCAAGATTTCCAAGGTACCTATCATGTTTCTTCTTCGAGAGA 400
AGGAGTACCTATAACCAACGGGAACAATTACCAATAGTGACCACTCTTTAGGCGTTCTAAAGGTTCCATGGATAGTACCAAGAAAGAGTCTCTCT

50 L M D I G L P L F N G Y H M C Q E I R K I S K V P I M F L S S R D 82

401 CCAGGCTATGGATATTGTCATGGCAATCAATATGGGGCGGATGACTTTGTGACCAAGCCTTTTACCAGCAGGTTCTTTAGCTAAGGTTCAAGGCTTG 500
GGTCCGATACCTATAACAGTACCTTAAGTTATACCCCGCTACTGAAACACTGGTTCGGAAAGTGGTCTGCCAAGAAATCGATTCCAAGTCCGAAAC

83 Q A M D I V M A I N M G A D D F V T K P F D Q O V L L A K V Q G L 115

501 TTGCGTCTTCTATGAGTTTGGGCGTATGAGAGTTTGTGGAATATGCTGGTGTATCTCAATACCAAAATCCATGGATTTACATTTCAAGGGCAAG 600
AACGCAGCAAGGATACCTCAACCCGCACTACTCTCAACGACCTTATACGACCACAATAGGAGTTATGGTTTAGGTACCTAAATGTAATAGTTCCCGTTC

116 L R R S Y E F G R D E S L L E Y A G V I L N T K S M D L H Y Q G Q V 149

601 TCTTGAATTTGACCAAGATGAATTCAGATTTTACGCGTGTATTGAGCATGCGAGCAACATCGTAGCAGTGCAGCCTGATGCGGAACTTTGGA 700
AGAACTTAACTGGTTCCTTAAAGGTCTAAATGCGCACAATAAATCGTACGTCCGTTGTAGCATCGTGCCTGCTGGACTACGCTTGAACCTT

150 L N L T K N E F C I L R V L F E H A G N I V A R D D L M R E L M N 182

701 CAGTGACTTTTCATTGATGATAATACCTCTCTGCAATGCGCTCGTTTCGTAAGGTTTGGAGGAGCAGGATTTGATAGCATTATCGAGACCAAG 800
GTCACTGAAAAAGTAATACTATTATGGGAGAGACAGTTACACGAGCAACGCAATTTTCAACCTCTCTGCTCCATACCATCTTAAATAGCTCTGCTTC

183 S D F F I D D N T L S V N V A R L R K K L E E Q G L V G P I E T K 215

801 AAAGGAATAGGGTACGGATTGAAGCATGCTTGATTGSAACAATTTTCTAGCCTATCTGCGCTCCCGTAGTCTCTTTTATCTATCTGCTTTCTTTG 900
TTTCCCTTATCCCATGCTAACTTCGTACGAATAACCTTTGTTAAAAAGATCGGATAGACGCGAGGSCATCAGCAGAAAAATAGATAGACGAAAGAAAC

216 K G I G Y G L K H A * 226

901 GCATTTCTGCTTACTCTTTCACTTTTATTTGCGAGTCTAGSAAATTTACTTCTCTAGTTTCTTCTTGTGTTGCTTTGTAAACCATCTTATTTTCA 1000
CGTAAAGAACAGATGAGAAAGTCAAAAATAAACGGTCAGATCTTAAATGAAGGAGATGAAAAAGAGAACACAAACGAAACATTTGGTAGAATAAAAGT

g9p1493

Fig. 5

(SEQ ID NO:14) 1 TAAAGACACTGGAACGACCAACACCTTCCGCATTTTAGGTAAAGAAAGCTGGTATGGCAACCTTTGTGATTGACTTTTCAAAGCAACCTAGCAACGCTG 100
(SEQ ID NO:15) 1 ATTTCTGTGACCTTGTGCTGTGTGGAAGGCGTAAATCCATTCTTTGACCATACCGTTGGAAACACTAACTGAAAAAGTTTCTTGGGATCGTTGCGAC
(SEQ ID NO:13) 1 K D T G T Y N T F R I L G K K A G M A T F V I D F F K O T L A T L 33

101 CTTCCGATTATTTTCACTCAAGGCGTTTCTCTCTCATCTTTGGACTTTTGGCTGTTATCGGCCATACCTTCCCTATCTTTCCAGGATTTAAAGGTG 200
GAAGGCTAATAAAAAGTAGATGTTCCGCAAGAGGAGAGTAGAAACCTGAAAACCGACAATAGCCGATATGCAAGGATAGAAACGTCTAAATTTCCAC
34 L P I I F B L Q G V S P L I F G L L A V I G H T F P I F A G F X G G 67

201 GTAAGGCTGTGCAACCACTCTCGAGTCAATTTCCGATTTCGCCCTATCTCTGTCTCTACCTTGGGATTATCTTCTTTGGACTCTCATATCTTGGCAG 300
CATTCGGACAGCGTTGGTCACGACCTCACTAAAGCCTAAACCGGATAGAAACAGAGATGGAAACCTAATAGAAGAAACCTGAGAGTATAGAACCTTC
68 K A V A T S A G V I F G F A P I F C L Y L A I I F F G L S Y L G S 100

301 TATGATTTCACTGTCTAGTGTCAAGCATCGATCGCGGCTGTTA 344
ATACTAAAGTGACAGATCACAGTGTCTAGCTAGCGCGCAAT
101 M I S L S S V T A S I A A V 114

gcp1507

Fig. 6

(SEQ ID NO:17) 1 CTAAGGTAATGAATGAAAGTATAAAATTAATGCTCTATCTTACATGGGAATTCGTCTTCAATATTTTCCATCCTAACTGGAACCTATG 100
(SEQ ID NO:18) GATTTCCATTTAACTTACTTTTCATATTTTAACTTACGAGATAGAAATGTACCCCTTAAGCACAGAACTTATAATAAAAAGGGTAGGATTGACCTTGGATAC
(SEQ ID NO:16) 1 M K S I K L N A L S Y M G I R V L N I I F P I L T G T Y V 29

101 TCGCGCGTGTCTTGGACCGAAGTCACTATGGTTACTTCAACTCAGTCGACACTATTTGTCTTTCTTGGCCCTTGGCACTTATCGTGTCTATAACTA 200
AGCGCGCACAGAACTCGCTTGACTGATACCAATGAAGTTGAATCAGCTGTGATAAAACAGTAAAAGAACGGGAAACGTTGAATACCAAGATATTGAT
30 A R V L D R T D Y G Y P N S V D T I L S F F L P F A T Y G V Y N Y 62

201 CGGTTTAAGGGCTATCAGTAATGTCAAGGATAACAAAAAGATCTTAACAGAACCTTTCTAGTCTTTTATTGTCATCGCTTGTACGATTTTGACC 300
GCCAAATTCGGATAGTCATTACAGTTCTTATTGTTTTCTAGAAATGTCTTGGAAAAGATCAGAAAAAATAAACACGTAAGCGAACATGCTAAAACCTGG
63 C L R A I S N V K D N K K D L N R T F S S L F Y L C I A C T I L T 95

301 ACTGCTGTCTATATCCTAGCCTATCCTCTCTTCTTACTGATAATCCAATCGTCAAAAAGGTCTACCTTGTATGGGGATTCAACTCATTGCCAGATT 400
TGACGACAGATATAGGATCGGATAGGAGAGAAGAAATGACTATTAGGTTAGCAGTTTTTCCAGATGGAAACAATACCCCTAAGTTGAGTAAAGGCTCTAAA
96 T A V Y I L A Y P L F F T D N P I V K K V Y L V M G I Q L I A Q I F 129

401 TTTCATCGAATGGGTCAATGAAGCTCTGAAAAATTACAGTTTCTCTTTACAAAACCTGC 460
AAAGTTAGCTTACCCAGTTACTTCGAGACCTTTAAATGTCAAGAGAAAAATGTTTTGACG
130 S I E W V N E A L E N Y S F S F T K L 148

gcp1311

Fig. 7

(SEQ ID NO: 20) 1 CCTCCATTACCGTGCATGCAATTCACGTATGTAATGATTTTATGGACAACTGGAGAGCAGGACGAGGAAATGTATGTTTGTGACGAGTTGCTATACA 100
(SEQ ID NO: 21) CGACGGTAAATGGCACTACCTAAAGTGCATACATTACTAAAAATACCTGTTGCAGCTCTGTCCTGCTCTTACATACAAAACACTGCTCAACGATATGT

101 GGGAGTAGGCATGCAGATTCAAAAAGTTTAAAGGGCAGTCTCCCTATGGCAAGCTGTATCTAGTGGCAACGCCGATTGGCAATCTAGATGATGACT 200
CCCTCATCGGTACGCTAAGTTTCTTCAAAATTCCTCCGTACAGGGGATACCGTTGCACATAGATCACCGTTGGCGCTAACCGTTAGATCTACTATACTGA

(SEQ ID NO: 19) 1 M Q I O K S F K G O S P Y G K L Y L V A T P I G N L D D M T 30

201 TTCTGTCTATCCAGACCTTGAAGAAGTGGACTGGATTGCTGCTGAGGATACGGCAATACAGGGCTTTTGTCTCAAGCATTTTGACATTTCCACCAAGC 300
AAAGCACGATAGGTCTGGAACCTTCTTCACTGACCTAACGAGGACTCTATGCGCGTTATGTCCGAAACGAGTTCTGTAAGTGTAAAGGTGGTTGG

31 F R A I Q T L K E V D W I A A E D T R N T G L L L K H F D I S T K Q 64

301 AGATCAGTTTTCATGAGCACAATGCAAGGAAAAAATTCCTGATTTCATTGCTTGAAGCAGGGCAAGTATTGCTCAGGTCTCTGATGCCGGTTT 400
TCTAGTCAAAAGTACTCTGTGTACGTTTCTTTTAAAGGCTAAACTAACCAAGAACTTCTGTCGCTTTCATAACGAGTCCAGGACTACGGCCAAA

65 I S F H E H N A K E K I P D L I G F L K A G O S I A Q V S D A G L 97

401 GCCTAGCATTTTCAAGCCCTGGTCAATGATTAGTTAAGGCAGCTATTGAGGAAGAAATTCAGTTGTGACTGTTCAGGTACCTCTGCAGGAATTTCTGCC 500
CGGATCGTAAAGTCTGGGACAGTACTAAATCAATTCCTGCGATAACTCTCTTTAACGTCAACACTGACAGGTTCCATGGAGAGCTCTCTAAAGAGCGG

98 P S I S D P G H D L V K A A I E E E I A V V T V P G T S A G I S A 130

501 TTGATTGCCAGTGGTTTAGCGCCACAGCCACATATCTTTACGGTTTTTACCGAGAAATTCAGCTCAACAGAAGCAATTTTTCGCTCTAAAAAGATT 600
AACTAACCGTCAACCAATCGCGGTGTCGCTGTATAGAAATGCAAAAAATGGCTCTTTAGTCCAGTTGTCTTCTGTAAGAAACCGAGATTTTCTCTAA

132 L : A S G L A P O P H I F Y G F L P R K S G Q O K O F F G S K K D Y 164

601 ATCTGAAACACAGATTTTATGAATCAGCTCATCGTGTAGCAGACCGTGGAAAAATATGTTAGAAGTCTACGGTGACCGCTCGGTGTTTGTGTCAG 700
TAGGACTTTGTGCTAAAAAATACTTAGTGGAGTAGCACATCGTGTGTCAACCTTTATACAATCTTCAGATGCCACTGGCGAGCCAAACAAACAGTGC

165 P E T Q I F Y E S P H R V A D T L E N M L E V Y G D R S V V L V R 197

701 GGAATTGACCAAAATCTATGAAGAAATACCAAGAGGTACAAATTTCTGAATTCCTGGAAAGCATCTCTGAAACGTCTCTCAAGGGTGAATGTCTTCTGATT 800
CTTTAACTGGTTTATGATCTTCTTATGGTTTCTCATGTTAAAGACTTAACGACCTTTCTGAGAGACTTTGCAGAGAGTTCCACTTACAGAGAGCTAA

198 E L T K I Y E E Y O R G T : S E L L E S I S E T S L K G E C L L I 230

801 GTTGAAGGTGCCAGCAAGGTGTGAGGAAAAGGATGAGGAAGACTTGTCTTAGAAATCCAAGCCCGTATCCAGCAAGGCATGAAGAAAAATCAAGCTA 900
CAACTTCCACGGTGGTTCCACACCTCTCTTCTACTCTCTCTGAACAGAACTTTTAGGTTCCGGCATAGGTGTTCCGTACTCTCTTCTAGTTCCAT

231 V E G A S K G V E E K D E E D L F L E I Q A R I Q Q G M K K N Q A : 264

901 TTAAGGAAATAGCTAAGATTTACAGTGGAAATAGAGTCAACTCTACGCTGCTTACCAGACTGGGAAGAAAAACAATAAAGGAGACAGGATGTAATAA 1000
AATTCCTTTATGATTTCAATGGTCACTTATCTCAGTTGAGATGCCAGGATGGTGTGACCCCTCTTTTGTATTTCCTCTGTCTCATATTAT

265 K E I A K I Y O W N K S C L Y A A Y H D W E E K Q * 290

gcp1518

Fig. 8 (Sheet 1 of 2)

(SEQ ID NO: 23) 1 ATGGCTTGGTTAAAAAAGGTGGCAATGCTCTTAACTGCAAGTTATTGGCTGTAGCAATATAAATCTATTTCTACATATTTTAAACGTTCTACGAG 100
(SEQ ID NO: 24) TACCGAACCAATTTTTTCCACGTTTACGAGAAATTCAGCTTCAATAACGGACATGTTATTTAGATAAAGGATGTTAAAAAATTTGCAAGATGCTC

101 TTAAATTTGAAACGTTTAGCTTGTGGTATAATAGATTTATGGATAAAAAATATGAAAAAATCTCTCAGGATTTGGGAGTCACTTTAAAGCAAATTTGATACC 200
AATTAACCTTTGCAATCGAACACCATATTTATCTAAATAGCTATTTTATACCTTTTATAGAGAGTCTTAAACCTTCACTGCAATTTCTTTAACTATGG

(SEQ ID NO: 22) 1 M D K K Y E K I S Q D L G V T L K Q I D T 21

201 GTTCTAAGTTTGACAGCTGAAGGGGCGACTATTCCCTTTATCGCGCTTATCGCAAGGACATGACTGGTAGTCTGGATGAGGTGGCGATTAAAGCTATTAT 300
CAAGATTCAAACTGTGCACTTCCCGCTGATAAGGGAAATAGCGGCAATAGCGTTCTCTGTACTGACCATCAGACTTACTCCACCGCTAAATTCGATATAT

22 V L S L T A E G A T I P F I A R Y R K D M T G S L D E V A I K A I I 55

301 TTGATTTGGATAAAAGTCTGACAAATCTCAATGACCGTAAGCAAGCTGTCTAGCTAAGATTCAAGAACAGGTAAAGTGAACCAAGGAATGCAAGAACG 400
AACTAAACCTATTTTCAAGCTGTTTAGAGTTACTGGCATTCCTTCCAGACAGATCGATTTCTAAGTTCTTGTCCATTCACTGGTTCTTAACTCTTCTGG

56 D L D K S L T N L M D R K E A V L A K I O E O G K L T K E L E E A 88

401 TATCTTAGTTGCCGAAAAATTAGCAGACGTTGAAGAACTCTATCTTCTTATAAGGAAAGCGTGTACCAAGGCAACCAATGCCCGTGAAGCTGGACTC 500
ATAGAAATCAACGGCTTTTAACTGCTGCAACTCTTGAGATAGAAAGGAATATCTCTTTCGACGATGGTTCCGTTGGTAAACGGGCACTTCCAGCTTGAG

89 I L V A E K L A D V E E L Y L P Y K E K R R T K A T I A R E A G L 121

501 TTTCTCTTCTGCTGTTTGAATTTGCGAATATAGTTGACTTAGAGAAAGAAAGCTGAAAGTTCTCTGTGAAGGATTTGCACTGGCAAGCAAGCTTGA 600
AAAGGAGAACGAGCAAACTAAAGCTCTTATATCAACTGAATCTCTTCTTCCGACTTTCAAGCAGACACTTCTTAAACGCTGACCGTTCTTCCGAACT

122 F P L A R L I L Q N I V D L E K E A E K F V C E G F A T G K E A L T 155

601 CCGGTGCAGTTGATATTTGGTGAAGCCCTTATCGGAAGATGTGACCTTGGCTTCTATGACTTATCAGGAAGTGTGAGACACTCTAACTCACTTCTCA 700
GGCCAGCTCAACTATAAACCCAGCTTCCGAATAGCTTCTACACTGGAAAGCAAGATCTGAATAGTCTTCAAGCTTGTGAGATTGAGTGAAGAGT

156 G A V D I L V E A L S E D V T L R S M T Y O E V L R H S K L T S O 188

701 AGCCAAAGGATGAAGCTTTGATGAAAAGCAGGTTTTTCAATTTATTATGATTTTTTCAAGACAGTTGGAACTATGCAAGGCTATCTGACTTGGCTCTC 800
TGGCTTCTTACTTTCAGAACTACTTTTCTTCCAAAAGTCTAAATAAATACTAAAAAGTCTCTGTCAACCTTGATACGTTCCGATAGCATGGAACCGAGAG

189 A K D E S L D E K O V F O : Y Y D F S E T V G T M Q G Y R T L A L 221

801 AATCTGGGAGAACTTGGTGTCTTGAAGATCGGTTTTGAACATCGGACCGACGTTATTTGGCTTCTTTGGTACTCGTTTCAAGGTGAAAAATGCTT 900
TTAGCACCTCTCTTTGAACCAAGAACTTCTAGCCAAAATCTGACCTGCTGGCTGGCATAAGAACCGGAAGAACGATGAGCAAGTTCCACTTTTACGAA

222 N R G E K L G V L K I G F E H A T D R : L A F F A T R F X V K N A Y 255

901 ATATTGATGAAGTTGTTCAAGCAATCCGTTAAGAAAAAGGCTTGGCTGCTATTGAGCGTCTGATTCCGACAGAAATTAAGTGAAGAAAGCTGAAGAGGAGC 1000
TATAACTACTTCAACAACTGTTAGGCAATTTCTTTCCAGAACGGACGATAACTCGCAGCATAGCCCTGTTAATTGACTCTTTGACTTCTCCCTCG

256 : D E V V O O S V K K K V L P A I E R R I R T E L T E K A E E G A 288

1001 TATCCAACTTTTTCTGACAACTCGGCAATCTCTTCTGCTTCTTCTTCTTCACTGAAAGGGCGGCTGGTTCTTGGATTTGACCCAGCTTTTCTGACAGTGGC 1100
ATAGGTTGAAAAAGACTGTTAGACGCTTAGAGGAGAACCAACGAGTGACTTCCCGCGCACCAAGAACCTAACTGGTGGGAAAGCATGTTCCACGG

289 : Q L F S D N L R N L L L V A P L K G R V V L G F D P A F R T G A 321

1101 AAGTTAGCTGTGTTGATGCAACAGGAAAAATGCTGCAACTCAGGTTATTTATCTGTTTAAACCAAGCATCAGCTGTTCAATCGAAGAACCAAGAAAG 1200
TTCAATCGACAGCACCTACGTTGCTTTTACGACTGTTGAGTCCAATAAATAGGACAAATTTGGTGTAGTGGAGCAGTTTACGTTCTTGGGTTCTTTC

122 K L A V V D A T G K M L T T O V : Y P V K P A S A R Q I E E A K K D 355

1201 ATTTAGCAGATTTAATTTGGTCAATACGGGTAGAGATTTATGCAATTTGGAAATGGAACCGCCAGCTGTTGAAAGTGAAGCTTTTGTAGCCGAACTTCTGAA 1300
TAAATGCTTAAATTAACGAGTTATGCCACATCTTAATAACGGTAACCTTACCTTGGCGGTGAGCACTTTCACTTGGAAACATCGCTTCAAGACTT

356 L A D L I C C Y G V E : I A I G H G T A S R E S E A F V A E V L K 388

Fig. 8 (Sheet 2 of 2)

10/30

1301 AGATTTCCTGAAGTCAGCTATGTTATCGTTAATGAAGTGGTCTTCTGTCTATTCTGCCAGCGAACTTCTCTCAGGAGTTTCCAGACTTGACCGT... 1400
TCTAAAGGGACTTCAGTCGATACAAATAGCAATTACTTTCACCACGAAGACAGATAAGAAGGTCTTGAAAGAGCAGTCTCTCAAAGGTCTGAAGTGGCAA

389 D F P E V S Y V I V N E S G A S V Y S A S E L A R O E F P D L T V 421

1401 GAAAAACGCTCTGCCATTTCATCGCCCTCGTTTGCAGATCTCTTGCAGAAATGGTCAAAATCGATCCTAAGTCAATTGGTGTGGTCAATACCAAC 1500
CTTTTTCGAGACGGTAAGATAGCGGGCAGCAACGTTCTAGGAGAACGCTTAACCAAGTTTAACTAGGATTCAAGTTAACCAAGCCAGTTATGGTTG

422 E K R S A I S I A R R L Q D P L A E L V K I D P K S I G V G Q Y Q H 455

1501 ACCATGTCAGTCAGAGAACTATCTCAGACTCTGGACTTGTGTGATACAGTGGTTAACCAAGTGGTGTCAATGTCATACAGCTAGCCAGCTCT 1600
TGCTACAGTCAGTCTTCTTTGATAGACTCTCAGACCTGAACAACAGCTATGTCAACCAATGGTTCAACCAAGTTACAGTTATGTCGATCGGGTCGAGA

456 D V S Q K K L S E S L D F V V D T V V N Q V G V N V N T A S P A L 488

1601 TCTTTCACAGCTAGCTGGACTCAACAAAATATCTCTGAAAATATTGTCAAATACCGCGAGGAAGGAAGAAAAATCACTTCAAGCGCCCAATCAAGAAA 1700
AGAAAGTGTGCATCGACCTGAGTTGTTTGTATAGAGACTTTTATAACAGTTTATGGCGCTCCTTCTTCTTCTTTAGTGAAGTGGCGGGTTTATGTTCTT

489 L S H V A G L N K T I S E N I V K Y R E E E O K I T S R A Q I K K 521

1701 GTTCTCTGTCGGGAGCCAGGCTTTGAGCAGGCTGCTGTTTCTTCTGATCCCTGAAAGTAGCAATATCCTTGATAATACAGGAGTTCAACCAGAG 1799
CAAGGAGCAGACCTCGTTTCCGAAACTCGTCCGACGACCAAGGAAGCATAGGGACTTTCATGTTATAGGAATATTATGCTCTCAAGTGGGTCTC

522 V P R L G A K A F E Q A A G F L R I P E S S N I L D N T G V H P E 554

gcp1546

Fig. 9

11/30

(SEQ ID NO: 26) 1 TACTGGGGCAAGGTTTCTTACCCGTGTTCTGAATGTGAAGTCTTTCTTGAAAATGGTGAAGTTAAGATTTTCAGAGCACTCAACGAAGCCAGNATCCGC 100
(SEQ ID NO: 27) ATGACCCCGTTCCCAAAGAATGGGACAAGACTTACACTTCGAGAAAGAACTTTACCACTTCAATTCATAAAGTCTCOTGAGTTGCTTCGGTCHTAGGGC
(SEQ ID NO: 25) 1 T G A R V S Y P V L N V K V F L E N G E V X I F R A L N E A X I R 33

101 ACOTCTCATCGAACCATGCTCCAGATATTGTAAATGGTGTTCCTTTGAACGTTTTCTGGAGACGGGCTAACAGTTTCCACACCGACTGGTAGTA 200
TCCAGACTAGCTTGGTACCACCGTCTATAACAATTATTACCAGAGGGAACTTGCAAAAGCACCTCTGCCCGATTGTCAAGCTGTGGCTGACCATCAT
34 R S D R T M V A D I V I N G V P F E R F R G D G L T V S T P T G S T 67

201 CTGCTATAACAAGTCTCTTGGCGGTGCTGTTTACACCCCTACCATTTGAAGCTTTGCAATTAAACGGAGATTGCCAGCCTTAATAATCGTGTCTATCGAAC 300
GACGGATATTGTTTACAGAGAACCGCCACGACAAAATGTGGATGGTAACCTTCGAAACGTTAATTGGCTCTAACGGTCCGGAATTAATTAGCACAGATAGCTTG
68 A Y N K S L G C A V L H P T I E A L Q L T E I A S L N N R V Y R T 100

301 ATTGGGCTCTTCCATTATTGTGCTTAAGAAGCATAAGATTGAACCTTATTCCAACAGAAACGATTATCATCTATTTCGGTTGACAATAGCGTTTATTCT 400
TAAACCGAGAGGTAATAACACGGATTCTTCTATTCTAACTTGAATAAGGTGTTCTTTGCTAATAGTATGATAAAGCCAACTGTTATCGCAATAAGA
101 L G S S I : V P K R D K I E L I P T R N D Y H T I S V D N S V Y S 133

401 TTCGTAATATTGAGCGTATTGAGTATCAAAATCCACCATCATAGATTCACTTTGTCCGCACTCCTAGCCATACCACTTTCTGGAAACCGTGTAAAGGATG 500
AAGGCATTATAACTCGCATAACTCATAGTTTAGCTGGTAGTATTCTAAGTGAAACAGCGCTGAGGATCGGTATGGTCAAAGACCTTGGCAAAATTCCTAC
134 F R N I E R I E Y Q I D H H K I H F V A T P S H T S F W N R V K D A 167

501 CCTTTATCGGTGAGGTGGATGAATGAGGTTGAATTTATCGCAGATGAACATGTCAAGGTTAAGACCTTTTAAAAAA 578
GGAAATAGCCACTCCACCTACTTACTCCAACTTAAATAGCGTCTACTTGTACAGTTCCAATTCGGAAAAATTTTT
168 F I G E V D E * 175

gcp1551

Fig. 10

(SEQ ID NO: 29)¹ CGCTCTAAAAGAAACCTACTCGAGACTCATAGATGGGAAGTACTATTATTTGATCCTTTATCCCGAGAGATGGTTGTGGCTGGCAATATATACCTGCT 100
(SEQ ID NO: 30) CCGAGATTTTCTTTGGATGACCTCTCACTATCTACCTTCATGATAATAAACTAGGAATAGGCCTCTCTACCAACAGCCGACCTTATATATGAGCGA
(SEQ ID NO: 28)¹ M V V G W O Y I P A 10

101 CCACACAAGGGGTTACGATTGGTCCTTCTCCAAGNATAGAGATTGCTCTTAGACCAGATTGGTTTATTTTGGTCAAGATGGTCTTACAAGAATTG 200
GGTGTGTTCCCCCAATGCTAACCCAGGAAGAGGTTCTTATCTCTAACGAGAACTGGCTTAACCAAAATAAAACCACTTCTACCAAGAAATGTTCTTAAAC
11 P H K G V T I G P S P R I E I A L R P D W F Y F G O D G V L Q E F V 44

201 TTGGCAAGCAAGTTTATAGAAGCAAAAAGTCTGCTACGAATACCAACAAACATCATGGGGAAGATATGATAGCCAGCAGAGAAACGAGTCTATTATTTGA 300
AACCGTTGGTTCAAAATCTTCGTTTGGAGATGCTTATGGTTGTTTGTAGTACCCCTTCTTATACTATCGGTTTCGTTCTTTGCTCAGATAATAAACT
45 G K Q V L E A K T A T N T N K H G E E Y D S Q A E K R V Y Y F E 77

301 AGATCAGCGTAGTTATCATACTTTAAAACTGGTTGGATTATGAAGAGGGTTATTGGTATTATTTACAGAAGGATGGTGGCTTTGATTCTCGCATCAAC 400
TCTAGTCGCATCAATAGTATGAAATTTTGAACCACTAAATACTTCTCCCAATAACCATATAAATGTTCTCTACCAACCGAACTAAGAGCGTAGTTG
78 D Q R S Y H T L K T G W I Y E E G Y W Y Y L Q K D G G F D S R I N 110

401 AGATTGACGGTTGGAGAGCTAGCAGCGTGGTTGGTTAAGGATTACCTCTTACGTATGATGAAGAGAAGCTAAAAGCAGTCCATGGTACTATCTAGATC 500
TCTAACTGCCAACCTCTCGATCGTGACCAACCAATTCCTAATGGGAGAATGCATACTACTTCTCTCGATTTCGTCGAGGTACCATGATAGATCTAG
111 R L T V G E L A R G W V K D Y P L T Y D E E K L K A A P W Y Y L D P 144

501 CAGCACTGGCTGGCAAAACCTTGGCAACAAATCGTACTACCTCGCTTCATCAGGAGCTATGGTAACTGGCTGGTATCAAGATGGTTTAACTGGTACTA 600
GTCTTGACCGAGCGTTTGGAAACCTTCTTTACCATGATGGAGGCAAGTAGTCTCGATACCATGACCGACCATAGTCTTACCAAAATTGAACCATGAT
145 A T G W O N L G N K W Y Y L R S S G A M V T G W Y Q D G L T W Y Y 177

601 CCTAAATGCAGGTAATGGAGACATGAAGACAGGTTGGTTCCAAGTCAATGGTAACTGGTACTATGCCTATGATTCAGGTGCTTTAGCTCTTAATACCACA 700
GGAATTACGTCCATTACCTCTGTACTTCTGTCCAACCAAGGTTCAAGTACCATGACCATGATACGGATCTAAGTCCACGAAATCGACAAATTATGGTGT
178 L N A G N G D M K T G W F Q V N G N W Y Y A Y D S G A L A V N T T 210

701 GTAGGTGGTTACTACTTAAACTATAATGGTGAATGGTTAAGTAATGAAGGCTAATTTGTAACCTGTGATGGATCTTAACTTTGTATAATAGGTGGATAA 800
CATCCACCAATGATGAATTTGATATTACCACTTACCAATTCATTACTTCCGATTAACTTTGACACTACCTATGAATTGAAACATATTATCCACCTATT
211 V C G Y Y L N Y N G E W V K * 225

gsp1561

Fig. 11

(SEQ ID NO: 32) 1 TTTTATGGATATTTATATTAAAGAAAGCCATTATTACCAAGTTCAGTCCGATGATACCGAGCTGTTCTTAGCAGATAAGTTTCTCAATATTACTCCAAAA 100
(SEQ ID NO: 33) AAAATACCTATAAATAATAATTCTTTCCGTAAATAAGTGGTCAAGTCAGGCTACTATGGCTCGACAAGAAATCGTCTATTCAAAGAGTTATAATGAGGTTTT
(SEQ ID NO: 31) 1 M D I Y I X K A I I H O F S P D D T E L F L A D K F L N I T P K 32

101 ATCGAAGAATACCTACGTAAAAAAATGAACATGCTATTGAGATGAAGCCAGACTGGGATTTTCAAGAAAGAAATCCCTTCTTCAATCATATTACAG 200
TAGCTTTCTATGGATGCAATTTTAACTTGTACACATAAGTCTACTCGGTTCTGACCTAAAGGCTTCTTCTTTAGGGAAGAAAGTACTATAATGTC
33 I E E Y L R K K I E H V Y S D E A K T G I F E E E N P F F N H I T D 66

201 ACCGATTGTTGGACACATCAGTAACGCTGGCTAATCTCTGGAAGAGGAGTTTAGCATTCTGAAAAATCTCAAGACCAATGACTTGATTTTGTTCAAAT 300
TGCTAAACCAACCTCTGTAGTCATTGCGACCGATTAGAGACCTTCTCCTCAATCGTAAGAGCTTTAGAGTCTGGTTACTGAACATAAAACCAAGTTAA
67 D L L E T S V T L A N L W K E E F S I S E N L K T N D L I F V Q F 99

301 TTCTAAAGAAAGGTGTAGAACATTTGCTTTCTTGGCAATTGCTTGGGAGAGCTTCAACCCCTCGGAGGAGAAAGTTGATAATCAATCAAGCTGACT 400
AAGATTCTTCCACATCTTGTAAAGCGAAAGAACGCTTAAACGGACGCTCTGGAAGCTGGGTGGAGCTTCTCTTCACTATTAGTTAGTTGAGCTGA
100 S K E G V E H F A F L R I A L R E T L T H L G G E V D N P I K L T 132

401 CAGAATAACCTGCTGGATTGGAACTGGTCTGACGAGGCTTGGTGGTCAATCTTCAGAGTCGCAAGTATCAGCTGATTGAAAAACGAATCAAGTACA 500
GTCTTATGGACCGACCTAAACCTTCCGACGACTCTCCGGAACCAAGGATCTAAAGAGGATTTTATAGTAGTTCTTCACTCTTTTGTCTAGTTTCACT
133 Q N H L P G F G T C A D E A L V V N L O S R K Y H L I E K R I K Y N 166

501 ACGGAGCTTTTGAAGTATTTTCAAGTAATCTTCTGCTGCTCTTCAAGATTCTCTTAAAAATCTATCAAGGAACTCGAAAAACAGCCGAGAG 600
TGCCCTGAAAAAATGATAAAAAAGTCTATTAGAAAGACGACGAGGATCTAAAGAGGATTTTATAGTAGTTCTTCACTCTTTTGTCTGGTCTC
167 G T F L N Y F S D N L L A V A P K I S P K K S I K E L E K T A O R 199

601 AATGCTGAATCTTTTAAACACAGATGATTTCAATTTCAATCAAGGTCAATCAGCTATTTTCAACCACTAGAAGAAAGCAATGAATGTCACCTGAG 700
TTAAGGACTTAGAAAAATGTTCTACTAAAAAGTTAAAGTAGGTTCCAGTTTGTAGTCGATAAAGTTGTTGGATCTTCTTCTGTTACTTAACAGTGGACTC
200 : A E S F N T D D F O F O S K V K S A I F N N L E E S N E L S P E 232

701 AAATGGCTAATGACCTTTTGAACAATCTGACGGCTCGTTTGGCTTTATTGACCAAGTCAGAGAAGCCGTACCAAGCTCTTCAATTTGATGAAA 800
TTTAAACGATTACTGGAAAACTGTTTGTAGACTGCCGAGCAAACTCGAAATAACTGGTTCACTCTTCTGGCATGGTCTTGGACAAGTTAAACTACTTT
233 K L A N D L F D N N L T A R L S F : D Q V R E A V P E P V Q F D E I 266

801 TTSATGCCAGTCGCCAATTAAGAAAAATGAAAAACAAAAACTCTCTTATCAATGGAAATGAGCTCATGTTCCCAATAACGTCTATCAAGACGCCGA 900
AACTACGGTCAGCGGTTAATTTCTTAACTTTTGGTTTTGAGAGGAATAGTTTACCTTAACTCGAGTAGCAAGGTTATTGAGATAGTTCTGCGGCT
267 C A S R Q L K K F E N Q K L S L S N G I E L I V P N N V Y Q D A E 299

901 GTCTGTTGAGTTTATCCAAAACGAAATGGAACCTACTCTATCTTAAATCAAAAATATCGAGGATATCCAAAGTAAATAATGTTTAAACGAATTCGAAGAG 1000
CAGACAACTCAAAATAGGTTTGTCTTTACCTTGGATGAGATAGAAATAGTTTATAGCTCTATAGGTTTCATTTATTACAAATTTGCTTAAAGCTTCTC
300 S V E F : Q N E N G T Y S I L : K N I E D I O S K * 1025

1001 TGCTGTACTAGCAGTCTCTCTTTTGTGGCTATAAGCTTACCGGTTTATCAAGATGTCAAAAGTCTATGACCTATCAACCCATGGTGGAGAAAT 1100
ACGAACATGATCTGTCAGAAAGAAAAACGACCGATATTTCGAATGGCGAAGTAGTTCTACAGTTTCTTCACTACTGATAGTTGGGTACCAAGCTCTTTA

gcp1580

Fig. 12

(SEQ ID NO: 35) 1 AAATGTGCTATAATACTAGAAAAATCTTGTGGAGGTTCCATTATGGCAATATTTTTCATGATTTTCTGATTGTTTGTGTGCTCTATTGGTGATGCT 100
(SEQ ID NO: 36) TTTACACGATATTATGATCTTTTATGAACACCTCCAAAGTAAATACCGTTATAAAAGTACTAAAAGACTAACAAACACACGAGGATAACCACTATCAG
(SEQ ID NO: 34) 1 M A I F F M I F L I V C V L L L V I V 19

101 AACTGAGTACAGTTTATGTCCTTCTCAGCAGTGGTGGCGATTATGGAACGCTTTGGCAAAATACCAAAAGGTTGCTAATAGCGGATTATCATATTCGCT 200
TGTGACTCATGTCAAATACACCAAGCAGTGTGAGCACCGCTAATAACTTGGCAACCCCTTATGGTTTTCACACGATTATCGGCATAAGTATAAGCGA
20 T L S T V Y V V R Q Q S V A I I E R F G K Y Q K V A N S G I R I R L 53

101 TGCCTTTTGGGATTGACTCGATTGCAGCACGGATTGAGTTGGCTTTGTCAAAAGTGATATTGTTGAGACTAAGACCAAGGACAATGTTTCGTTAT 300
ACGCAAAACCCCTAAGCTAAGCTGCTGCTAAGTCAACGGCAACAACTTTCACTATAACACCAACTCTGATTCTGGTTCTGTTACACAAGCAATA
54 P F G I D S I A A R I O L R L L Q S D I V V E T K T R D N V F V M 86

101 GATGAATGTAGCGACTCAGTACCGTGTCAACGAGCAGAGCTGCACAGATGCTTACTATAAATCATACGTCCAGAACTCTGAGTTAAATCTTATATCGAA 400
CATCTTACATCGCTGAGTCATGGCACAGTGTGCTGCTCGCACTGTCTACGAATGATATTGAGTATGAGGTCTTAGAGTCTAATTTAGATATAGCTT
87 M N V A T Q Y R V N E Q S V T D A Y Y K L I R P E S O I K S Y I E 119

101 GATGCTCTTCGCTCTTCTGTTCCAAAAATTAACCTTGGATGAATTGTTGAGAAAAAGATGAGATTGCCCTTGAGGTTCAACACCAAGTAGCAGAAGAAA 500
CTACGGAAGCGAGAAGACAGGTTTAAATGGAACTACTTAAACAACTCTTTTCTACTCTAACGGGAACCTCAAGTTGTGGTTCTGCTCTCTCTT
120 D A L R S S V P K L T L D E L F E R K K D E I A L E V O H O V A E E M 153

101 TGACCACTTACGGCTACATTATCGTGAAAACTTGATTACCAAGGTGAAACAGATGCAGAAAGTTAAGCAATCTATGAATGAAATCAATCGCGCGCAAG 600
ACTGGTGAATCGCGATGTAATAGCACTTTGGAACTAATGGTTCCAGCTTGGTCTACGCTCTCAATTCTGTAGATAGTACTTACTTTAGTTACGCGCGCTTC
154 T T Y G Y I I V K T L I T K V E P D A E V K Q S M N E I N A A O R 186

101 TAAGCGCGCTCGCAGCACAGAAATGGCGCAAGCTGACAGAAATTAATTTGTCAGTGCAGCTGAAGCGCAAGCAGAAAAAGACCGCTTATGGTGTGGGG 700
ATTGCGCCAGCGTCTGTTCTTAACCGCTTTCGACTGTTCTAATTTTAACAGTGACGTGCACTTCGGCTTCTGTTCTTTCTGGCGGAAGTACCAACCC
187 K R V A A Q E L A E A D K I K I V T A A E A E A E K D R L H G V G 219

101 ATTGCCCAACAAAGTAAAGCGATTGTTGGATGGATTGGCAGAGTCTATCAGCGAACTCAAGGAAGCAATGTTGGCATGACAGAAGAAATCATGTCTA 800
TAAAGGGTTGTTGATTCCGCTAACACCTACCTAACCGTCTCAGATAGTGGCTTGAGTTCTTGGTTACAAACCGTACTGTCTTCTGTTTATGACAGAT
220 I A Q O R K A I V D G L A E S I T E L K E A N V G M T E E Q I M S I 253

101 TCCCTTTGACCAACCACTATTTCGATACCTTGAATACCTTTCCTCTAAGGAAATCAAAACCATCTTTTACCAAAATCTCCAAATGGTGTGGATGATAT 900
AGGAGAACTGGTTGGTCATAAACCTATGGAACTTATGGAACGGAGATTTCCTTTAGTTTGGTAGAAAAATGGTTTATGAGGTTTACCAACCTACTATA
254 L L T N O Y L D T L N T F A S K G N Q T I F L P N T P N G V D D I 286

101 CCGTACACAAATCTTGTGAGCCCTTCGGCTGAGAAGAAATATAGACTAATACTCTTCGAAAAATCTCTTCAAACTACGTGAGCGTCTTTCGCGTATA 1000
GGCATGTGTTTAGAACAGTCCGGGAAGCGGACTCTCTTTATTATCTGATTATGAGAAGCTTTAGAGAAGTTTGATGAGTGGCAGCAGAACGGCATAT
287 P T Q I L S A L R A E K K 300

gsep1713

Pag. 13

(SEQ ID NO: 38) 1 CCTTCATATGGTGGATAAAATAGGGTTTATTTTGGAAACCTTTCCCTTTGTTTCAAAATGCTAAAAAANTGCTACAAATAGGAAAGCTTACTATTA 100
(SEQ ID NO: 39) GGAACATATACCACTATTTATCCCAAAANTAAAACTTTTGGAAAGGAAACAAAGTTTAAAGATTTTTHACCAATGTTATNTCCCTTTCGAATGATAAT

101 TCTGAATCAGCAGATTTCGAGAGAAAGGATTCATTTTGAATCAATAGGCTTTATGAAAAGCTGAAGGGGTTGTCTAGTAAAGAGCTGATTTTATGGG 200
AGACTTAGTCTGTAAACCTCTCTCTCTTAAGTAAAACTTTAGTTATCCGAAATAACTTTTCGACTTCCCAACAGATCATTTCTCGACTAAAAATAACCC
(SEQ ID NO: 37) 1 L K S I G P I E K L K G L S S K E L I L L G 22

201 AATTATCCTAAGTATCTTTTACCCCTTTTATCTTTTGTAGTTGTACTCTGTTTATATATTATCACTTTGATTTTACAGGAGACATGAAAAGTATCTTT 300
TTAATAGGATTCATAGAAAAATGGGAAATAGAAAAACATCAATGAGACAAATATATAATAGTCAAACTAAAAATGTCCTCTGTACTTTTCATAAGAA
23 I I L S I F L P F Y L F V V L C L Y I I S L I P T G D M K S I L 55

301 CAGAAAAATGGGGAGCATCCGATGCTCTCTCTCTTCTAGCTATAGTACTGTTTATATCCATCTTGCACAAAAATGGATGGGCTTGTGGCTTCAGTAG 400
GCTCTTTACCCCTCTAGGCTACGACGAGAAAAAGAAATCGATATCATGACAAATATAGGTAAGAAAGCTGTTTAACTACCCAGAACACCGAAGTCTAC
56 Q K M G E H P M L L L F L S Y S T V I S I L A O N W M G L V A S V G 89

401 GAATGTTCTATTACTATTTCTTTTGCCTATCATGCTGATTTATCCATAAAATCTTTCGATTGATTTTGCAGTTCTCTGTTTGGTAGTGTCTT 500
CTTACAAAGATATAATGATAAAGAAAAACCTGATAGTCACTAGCTAAAAATAGGGTATTTAAGAAAGCTAACTAAAAAGCTCAAGCAGAACCAACCATCACAGAA
90 M F L F T I F F L H Y O S I L S N K F F R L I L O F V L F G S V L 122

501 GTCAGCTGCTTTTCCAGTTAGAACATTTCCAAATGTCGAAGAAATTAACATATGCTTTTCTTTACCCCAATATGCAGGTGTCGATCAGAACCCGGCA 600
CAGTCGAGCAAAACCGTCAATCTTGTAAAGGTTTAACTCTCTTTAAATGATACGAAAAGAAAGTGGGTTATAGCTCCACACCGTAGTCTGGCCCGT
123 S A A F A S L E H F G I V K K F N Y A F L S P N H Q V W H Q N R A 155

601 GAAGTGACCTTCTTAATCCTAATTATATGGAATTAATTTGTTGTTCTGTTATTATGATGCTTTCTATCTGTTTACAAAGCAAGTTGAATTTGGTTGA 700
CTTCACTGGAGAAATTAGGATTAATAATACCTTAATAAACAAACAAAGACATAATCTAACGAAGATAGACAAATGTTGCTGTTTCACTTAACCAACT
156 E V T F F N P N Y Y G I : C C F C I M I A F Y L F T T T K L N W L K 189

701 AAGTATTCTGTGATGATGAGGCTTTGTTAATCTCTTGGTGAAGTTTACTCAAAATCGAACTGCTCTTCTGCTATTATCGCTGGAGCAATATATCTA 800
TTCAATAGACACACTAACCTCCGAAACAAATAGAGAAACCAACTTGAATGAGTTTACCTTGACGGAAAGGACGATATAGCGACCTCGTTAATAGAT
190 V F C V I A G F V N L F G L N P T O N R T A F P A I I A G A I I Y 222

801 TCTCTTAGGACTATTAAAACTGGAAGGCTTTTGGCTTAGTATTGGGCTCTTCGCAATGCTTTGAGTTTCTCTTTCTAGTATTGGGAGTTTGA 900
AGAGAAATGCTGATAATTTTGACCTTCCGAAAAACCGAATCATAAACCCAGAGCGCTAACCAACTCAAGGAGAAAGATCACTAAACCTCAAGCT
223 L F T T I K N W K A F W L S I G V F A : G L S F L F S S D L G V R 255

901 ATGGGTACTTTAGACTCTTTATGGAAGAACCCATTTCTATCTGGGATGCTGGATGCTCTTGTAAAGCAAAATCTTTTGGGTTGAAGGGCCATTGA 1000
TACCCATGAAATCTGAGAGATACCTTTCTGCTAAAGATAGACCTACGACCTACCGGAACAAATCTTTTAGGAAAAACCCCACTTCCCGGTAACT
256 M G T L D S S H E E R : S I W D A G H A L F K O N P F W G E G P L T 289

1001 CCTATATGCACTCTTATCTCGGATACATGCTCTTATCATGAACATGCCACAGCTTTTATATTGATACGATCTGAGTTACCGAAATTTGGGTACCAT 1100
GGATATACGTGAGAAATAGGAGCCTATGTACGAGGAATAGTACTTGTACGGGTGTCAGAAATATAACTATGCTAAGACTCAATGCTTAAACCCATGGTA
290 Y M H S Y P R I H A P Y H E H A H S L Y I D T I L S Y G I V G T I 322

1101 TTTATTAGTTTGTCTCTGTTGCTCTCTCTCTGATGATGATATGAGTCAGGAGTCGGGGAACGTCGATTATCGGCTTTATCTATCTTTCTCT 1200
AAATAATCAAAACAGAGACAAAGGAGCAAGCACTACTACTATCTAGTCTCTCAGCCCTTTGCAAGCTAAATAGCCGAAATAGATAGAAAGGAA
323 L L V L S S V A P V R L M H D M S O E S G K R P I I G L Y L S F L 355

1201 ACAGTGGTGGTGTGCAAGGAAATTTGACTTGGCTCTCTTGGATTGAGTCAGGCTTTATTTCTTGTACTATGTCAGTATTCAGTATTCAGTGGCTTTA 1299
TGTCAACCAAGACAGTGGCTTAAAACTGAACCGAGAGAGACCTAAGTCAGTCCGAAATAAAAAGACGATCAATACAGCTCATAGGTAAACCGAAAT
356 T V V A V H G : F D L A L F W I O S G F : F L L V M C S I P L A L 388

gcp222

Fig. 14

(SEQ ID NO: 41) 1 AAGGAGTGAACATCTGGCTCGTACTTCAATTGATGAAGATATGCGTCAATTTCTGTAAACAGTTGTCCGAACGGGTGTTCTGTCAAGACCCGGTAGA 100
(SEQ ID NO: 42) TTCTCACTTGTAGACCGAGCCATGAAGTTAACTACTTTTCATACCGCACTACTTAAAGCACATTGTCAACAGCGTTGCCACAAGCAGTTCTGGCGCATCT

101 AAAGGTTGTGGCTCCACAAGCTAGATCTGCTACTAACTACCGTGACAGCTGAACCCAGCTCATTCACATGGCTTTGATCGTCAATTTTATATGGCAGAA 200
TTTCCAACACCGAGGTGTTGATCTAGACGATGATTGATGGCACTCTGTCACTTTGGTCAGTAAGTGTACCGAACTAGCAGTAAAACTATACCGCTCT

201 ACAGTTGAATTCGCAAAACAAAATCCACGTCCTTTGGAACCAACTCAGGCACTGCTTTTGGTGATTGGGATCTTCCCGGTGAATCGATTGTTCTGTACAA 300
TGTCAACTTAAACGGTTTGTGTTTGGTGCAGCAAACTTGGTTGAGTCCGTAGACGAAACCACTAAACCTAGAGCGGCACCTAGCTAAACGATGTT

301 CAGATTGAGTCTTTCTCCAGTCCAGCGCTTTGAAGCCCAATTTCAAGATGAAGATGAATTCGATACACCTCCATTTTCAAAAAATCGTTAAGTAA 400
GTCTAAGTCAGCAAGAGGTGAGCTCCGCAAACTTCGGGTTAAAGTGTCTACTTCTACTTAACTATGTCGAGGTAAAGGTTTTTAGCAATTCAATT

(SEQ ID NO: 40) 1 M 1

401 TGAATGTAAAAGAAAATACAGAACTTGTCTTTCGAGAAGTTGCAGAGGCTAGTCTGAGTGCTCATCGAGAGAGTGCTTCGGTCTCTGTCAATTCAGTTAT 500
ACTTACATTTCTTTTATGCTCTTGAACAAAAGCTCTTCAACGTCTCCGATCAGACTCAGAGTAGCTCTCTCACCAAGCCAGAGACAGTAAAGTCAATA

2 N V K E N T E L V F R E V A E A S L S A H R E S G S V S V I A V : 14

501 CAAGTATGTAGATGTACCGACAGCGGAAGCTTTGCTTCCGCTAGGTGTTTATCATATCGGTGAAAATCGTGTAGATAAGTTTTCGAAAAAATATGAAGCT 600
GTTCATATCATCTACATGGCTGTGGCTTTCGGAACGAAGGCGATCCACAAGTAGTATAGCCACTTTTAGCACATCTATTCAAAGACCTTTTATACCTTCGA

35 K Y V D V P T A E A L L P L G V H H I G E N R V D K F L E K Y E A 67

601 TTAAGATCGAGATGTGACTTGGCAATTTGATTGGTACCTTGCAGAGCTAAGGTGAAGATGTCAATCAACGTTGATTATTTTCATGCAATTCGACT 700
AATTTTCTAGCTCTACACTGAACCGTAACTAACCATGGAACTTTCTGCACTTCCACTTTCTACAGTAAGTTATGCAACTAATAAGGTACGTAACTGTA

68 L K D R D V T W H L : G T L O R R K V K D V I Q Y V D Y F H A L D S 101

701 CAGTAAAGCTAGCAGGGGAAATTCAAAAAGAACTGACCGAGTCACTCAAGTGTCTCTTCAAGTAAATATTTCTAAAGAAAGAAAGCAACACCGTTTTC 800
GTCAATTCGATCGTCTCTTAAAGTTTTTCTTCACTGGCTCAGTASTTACAAAGGAAGTTCAATTTATAAGATTTCTTCTTCTGTTTGTGCAAAAAAG

100 V K L A G E : Q K R S D R V I K C F L C V N : S K E E S K H G F S 134

801 GAGAGGAACTGCTGAAAATCTTCCAGAGTTAGCCAGACTAGATAAGATTGAATATGTTGTTTAAAGACGATGGCACCCTTTGAGGGTAGCAGTGAG 900
CTCTCTCTTACGACCTTTAGAACGGTCTCAATCGGTCTGATCTATTCTAACTTATACAAACAAATTAAGTCTACCGTGGAAAACTCCGATCGTCACTC

135 R E E L L E : L P E L A R L D K : E Y V G L M T H A P P E A S S E 167

901 CAGTTGAAGAGATTTTCAAGCGGGCCCAAGATTTACAAAGAGAAATTCAGAGAAACAAATTCCAAATATGCTTTAGAGCACTGGCGGCGGTAC 999
GTCAACTTTCTTAAAGTTCCGCGGGTTCTAAATGTTTCTCTTTAAGTTCTCTTCTTTAAGTTTATACGGAATCTCTGTGACCGCGGCAATG

168 C L K E : F K A A C C L O R E : Q E R C : P N M P L E H T G G R Y 200

Pg. 15

(SEQ ID NO: 44)	1	GTACTCCAGTCCACTTTTAGCAGTAAGTTTATTATTCTTTTAAATCAGCCACAATTTCTGTCTTGAATCAGATTTTGGTAGGTAAGTTTGGTAATCTT	100
(SEQ ID NO: 45)		CATGAGGGTCAGGTGAAAAATCGTCACTCAAAATAATAAATGAAATTAAGTCGGTGTTAAAGAACAGAACTTACTCTAAAACCATCATCAAAACCAATTAGA	
(SEQ ID NO: 43)	1	T P S P L L A V S L L F T F N Q P Q F L V L N Q I L V G S L V I L	33
	101	ACTTATTGCATATATAGTTGTAAAAATCCCATTTTCTTATAGAAATGCTAGCTGCTATTTTTATTAGTGTTGATGATGAGATGGAAGATGCCACCAAGAGT	200
	34	L I A Y I V V K I P F S Y R M V R A I L F S V D D E M E D A A R S	66
	201	ATGGTGCTTCACTTTTATACATATGATGAAGGTTATCATTCATTATTTTACCGGTTGTTCTCTCTGTTATTGCTTTAACTTTAACTCTTTATTAA	300
	67	TACCCAGCAAGTGAAAAATATGATACTACTTCCAATAGTAAGGTAATAAATGCGCAACAGAGAGACAATTAACGAAATTTGAAATTCAGAAAAATAT	300
		H G A S P F Y T M K R V I I P F I L P V V L S V I A L N F N S L L T	300
	301	CTGACTTCGACTTATCTGTATTCCCTTTACCATCCCCCTAGCTCAACCATTAAGGTATTACCAATCCATCTGCAAGTGATGAACAGCAACATCTAATGCA	400
	101	GACTGAAGCTGAATAGACATAAGGAAATGGTAGGGGATCGAGTTGGTAATCCATAATGCTAAGCTAGACGCTCACTACTTTGTGTTGTAGATTACGTGT	133
		D F D L S V F L Y H P L A O P L G I T I R S A G D E T A T S M A Q	
	401	AGCTCTGGTATTGTTTATACAAATGTTCTGATGATTATTTCTGGAAACGGTATTATATCTTCAACAAGACCGGGGGCGTAAAGTAAGGAAATAATCATGA	500
	134	TGGAGACCAATAAACAAATATGTTAAACAGACTACTAATAAAGACCTTGCCATAAATGGAAGTGTTGTTCTGGCCCCGATTTCACTCTCTTTATTAGTACT	500
		A L V F V Y T : V L M I : S G T V L Y F T Q R P G R K V R K *	164
	501	CAGCCACTAGTCTTGGGTATCAAAATATTGAAATAGTTGTGAGGATTGTTTATCAGTAGTCAATGCTAGTATAATGCTTTAGACAGACGGAGCAAAATC	600
		GTGGTGATCAGAACCAATAGTTTATACTTTATCAACAGTCTTAAACAAATAGTCAATCAGTAACCATCATATTAACCAAAATCTCTCTCCCTCGTTAG	
	601	CCAGCCTCAGGCATCCGAACTTATAGTATTGTTGCTAGCTGCATGTTGATTATGATGACGAATGAATACGTTATCTTATAAATTTGGGACAGGAGAT	700
		GGTGGGACGTGGTAGGCTTGAATATCATAAACAAAGATCGAGCTACAACTAATAGTACTGCTTACTTATGCATAGAATATTTAAACCTGTCTCTCTA	
	701	CCTACAGGATTAGGAGCTCAAGTTATATCAGGTGTGGGTTTCTAGGCGTGGAAACGATTCTTATTACAGATAAAAAAGAAAAATACAGGCTGACAACTG	800
		GGATGTGCTAATCCTCGAGTTCAATATAGTCCACACCCAAAGATCCGCGACCTGCTAAGAATAATGTCTATTTTATATGTCGAGCTGTTGAC	
	801	CAGCAGGCATTTGGGCTTCCGCAGCAATGGATTAGCTATTGGAGTAGGTTTTTATCAGGCGAGCTCTTTTAGTAGGCCATTTCTGTTGGGGTGATATC	900
		GTGTCGTGTAACCCGAGAGCGTCTTAACTTAATCGATAACCTCATCCAAATAATCTCCCTCGAGAAATCATCGGTAAAGACAAACCCCACTATAG	
	901	CATGTTCCAAACCACTAAAAAATATCTGCAAAATCGTTCTAAAAATGATTGAATTTGATATAGTAGTTAAATCCTTTAG	978
		GTACAGGTTGGTGATTTTATAGACGTTTAGCAAGATTTACTAACTTAACATATATCATCAATTTAGGAAATC	

gdp273

Fig. 16

(SEQ ID NO: 47) 1 CAATGTGTTCCCGAACTTTTACAAAACATCTCTGAAAAAGAGTTTCAACACTCAAGACCAATTTGGTCAAAATAGGATGGTGTGTTGATGATG 100
(SEQ ID NO: 48) 1 GTTACACAAGGGCTTGAAAAATCTTTGTAGAAAGACTTTTTCTCAAGCTTGTGAGTTCTGGTTAAACCACTTTATCTACCAACACCAACTACTAC 100
(SEQ ID NO: 46) 1 M M 2

101 CACAGGATTAGACAAGAGTTGAAAAAGGGTGGAGCTGTCTCTACCTACAGAGACTGTTTATGGTCTTTTTTCCAAGGCTTAGATGAAAAAGCAGTTG 200
CTGTCTTAATCTGTTCTCAACCTTTCCACCTCGACAGCAAGATGGATGTCTCTGACAAATACCAGAAAAAGGTTCCGGAATCTACTTTTTCTCAAC 200

3 D R I R Q E L E K G G A V V L P T E T V Y G L F S K A L D E K A V D 36

201 ACCATGTTTACCAACTCAAGCTGTCTAGAGATAAGGCACTCAATCTCAATATCGGCTCTTTCAGGACATCTTGCACTTTTCAAAGAAATCAGCCAGC 300
TGTACAAATGGTTGAGTTTGCAGCAGGATCTCTATTCCGTGAGTTAGAGTTATAGCGGAGAAAGCTCTGTAGAACGTGAAAAGTTTCTTAGTGGTCC 300

37 H V Y Q L K R R P R D K A L N L N I A S P E D I L H F S K N O P A 69

101 TTATCTACAAAACTTGTAGAGACCTTTTGGCAGGTCCCTTGACCATTTCTCGAAGCCAATGACCGAGTTCCCTATTGGTAAATCTGACCTTGCA 400
AATAGATGTTTGTGAACATCTCTGAAAAAGGGTCCAGGAACTGGTAATAGAGCTTCGGTTACTGGCTCAAGGATAACCCATTAAAGATCGAAGCT 400

70 Y L O K L V E T F L P G P L T I I L E A N D R V P Y W V N S D L A 102

401 ACTATTGGATTTGGATGCCAGTCACTATGACACTGATTTAATTCGAGACAGAGTCCCTTGATTGGGCGGTCTGCCAATATCTCAGGTGAGGCA 500
TGATAACCTAAAGCTTACGGGTGAGTGGATAGTGTGACCTAAATTAAGCTCTCTGTCCAGGAACTAACCCGGCAGACGGTTATAGAGTCCAGTCCGTT 500

103 T I G F R M P S H P I T L D L I R E T G P L I G P S A N I S G Q A S 136

501 GTGGTGAACCTTTGAACAAATTTGAAAGATTTTACCAAGAGGTTCTGGGTCTGGAAGAGATGCTTTTCTAACTGGACAGGATTCACATTTGTTGGA 600
CACCATTTGGAACCTTTGTTAAGACTTCTAAATGTTTCTCAAGACCCAGAGCTTCTGCTAGGAAAAAGATTGACCTGTCTTAAGTTGATAACACCT 600

137 G V T F E Q I L K D F D Q E V L G L E D D A F L T G Q D S T I V D 169

601 TTTGCTCGAGACAGGTCAAAACTTACCAAGCGCAATTAACGAGAGATTTCTGCTCGGTGCCAGAGATTTCTTTGAGGAGGCTTGAATG 700
AAACAGACCTCTGTTCCACTTTAGAAATGGGTTCCGGTTAATTTGCTCTTCTAAGAAAGAGGCAACGGTCTCTAAGAAAACTCTCCGAACTTTAC 700

170 L S G D K V K I L P K A Q L N E K I F L L G C Q R F L L R R L E M 202

701 CTAAAGAGATTTGCAAGAAACAGATGTGAAGCGATATGTGACATCAACCAAGAGGCTTTGGGTTATCTTTAGTCCAGAGGAAACGGGTAGCCAACTAG 800
GATTTCTAAACGTTCTTTGTCTACACTTTGGCTATACACTGTAGTTGGTCTCCGAAACCAATATGAAAAATCAGGTCTCTTTGCGGATCGGTTGATC 800

203 L R D L Q E T D V K A : C D I N Q E A L G Y T F S P E E T A S O L A 236

801 CTAGACTGTCTCAGGATTECCATCTTTCTACTTGGTATGAGGATGCACTAATCATGTCTTACTTGGATATGTCCAGCTGAAGTTTACGAATCACT 900
GATCTGACAGAGTCTTAAGGCTAGTAAAGGATGAACCGATCTCTACGTCGATTAGTACAGAAAGAACTATACAGGTGCCACTTCAATGCTTAGTGA 900

237 R L S Q D S H H F L L C Y E D A A N H V L L G Y V H A E V Y E S L 269

901 CTATTCCAAGCAGGATTTAATATCTTAGCTTTAGCAGTTTCACTCAAGCGCAAGGTCAAGGTATCGGTAAGTTTACTACAGGTTGGAACAGAA 1000
GATAAGTTTCTGCTCTAAATTAAGAAATCGAAATCGTCAAGTGGAGTTCCGTTCCAGTTCCATAGCCATTTTCAATGATGTTCCCAACCTTGTCTT 1000

270 Y S K A G F N I L A L A V S P O A O G Q G I G K S L L O G L E O E 302

1001 GCCAAAAGATGTGGTTATGGGTTTATCCGTTTAAATCTGCCAATCATGCTCTGGGTGCTCATGCAATTTATGAAAAAGTTGGCTATATCTGTGATAAA 1100
CGGTTTTCTACACCAATACCCAAATAGGCGAATTTAAGACGGTTAGTAGGAGACCCAGGATAGTAAAAATCTTTTCAACCGATATGAACACTATTTT 1100

303 A K R C G Y C F : R L N S A M H R L G A H A F Y E K V G Y T C D K M 336

1101 TGCAGAAAGGTTTATTCGCATCTTTAGTTTGAATTTCTTATTTGTAATCAAACTAATGGACTAGTCACAATAAAGGAGAGAGCTATGATTTTGT 1200
ACGCTTTTGCAGAAATAGCGTAGAAATCAAACTAAAGAAATACATTTTAGTTTGAATACCTGATCAGTGTGTTATTTCTCTTCTGGATCTAAAAAC 1200

337 C K R F : R I F * 345

gcp206

Fig. 17 (Sheet 1 of 2)

(SEQ ID NO: 50) : AAGATAATAGAAAAAGAAATGTAACGAATGAGAGAAAAATGCGATTTGGAGATAATGGAATCGTAAAAAACTATGTTTGAGAAAAATACCTTGTATT
(SEQ ID NO: 51) : TTCTATTATCTTTATCTTACATTGCTTACTCTCTTTTACCGTAAACCTCTATTACCTTAGCATTTTGTATACAACTCTTTATTGGAAACAAATA 100

101 CCGTATTATCATGCTAGTAGCAAGTTTATGGGAATTTTTCGCACTGCAATTTGGTGCTTCACTAATCTATAAAATGATTCAAGAAAAATTTAGTGACTG 200
GCACATAAGTAGCATCATGCTTCAAATAACCTTAAAAACGTTGACGTTAACCGGAAGTCATTAGATATTTAACTAAGTCTTTTAAATCACTGAC

201 GGATTTCACGCCCTTTTAAAGTGAGAAGAAATAATGAGTATGTTTATAGATACAGCTAAGATTAAAGTCAAGGCTGTAATGGTGGCGATGGTATGG 300
CCTAAAGGGTCGGGAANAAATTTCACTCTTCTTTATTACTCATACAAAAATCTATGTCGATTCTAATTCAGTTTCGAGCATTACACCGCTACCATACC

(SEQ ID NO: 49) : M F L D T A K I X V K A G N G G D G M V 20

301 TTGCTTTTCGTCGTGAAAAATATGTCCTTAATGCGAGCCCTTGGCGTGGTATGCTGCTGTCGAGCCAAATGGTCTTCTGTTGACAGCAAGGACTACG 400
AACGGAAGCAGCACTTTTATACAGGGATTACCTTCGGGAACCCCACTACTACCAGCAGCCTCGTTACACAGGAAGCAACTGCTCTCTGATGC

21 A F R R E K Y V P N G G P W G G D G G R G G N V V F V V D E G L R 53

401 TACCTTGATGGATTTCCGCTACAATCGTCATTTCAAGGCTGATTCTGGTGAAGGGGATGACCAAGGGATGCAATGGTCTGGTGTCTGAGCACTTGA 500
ATGGAACTACCTAAAGGCGATGTTAGCAGTAAAGTTCCGACTAAGACCACTTTTCCCTACTGTTTCCCTACGTAACGACCAAGCACTCTGGAATCT

54 T L M D F R Y N R H F X A D S G E K G M T K G M H G R G A E D L R 86

501 GTTCGAGTACCAAGGTAAGGTAAGTCTGTCGATGCGGAGACTGGCAAGGTTTAAACAGATTTGATTGAACATGGGCAAGAAATTTATCGTTGCCACCGTG 600
CAAGCTCATGTTGTTCCATGCTGACAAAGCACTACGCTCTGACCGTTCCAAAAATGTTCTAACTAACTTGTACCCGTTCTTAAATAGCAACGGGTGCCAC

87 V R V P Q G T T V R D A E T G K V L T D L I E H G O E F I V A H G G 120

601 GTCTGGTGGACGTGAAATATTCGTTTCGGACACCAAAAAATCTGCACCGGAAATCTCGAAAAATGGAGAACAGGTCAGGAACTGAGTTACAAT 700
CAGCACCACTGCACCTTTATAAGCAAGCGCTGTGCTTTTAGGAGCTGGCCTTTAGAGACTTTTACCTCTGGTCCAGTCTGCACTCAATGTTAA

121 R G G R G N I R F A T P K N P A P E I S E N G E P G O E R E L O L 153

701 GGAATCAAAATCTTGGCAGATGTCGTTTAGTAGGATTCCCATCTGTAGGGAAGTCAACACTTTTAAAGTGTATTACCTCAGCTAAGGCTAAAAATGCT 800
CCTTGATTTTAGAACCGTCTACAGCCAAATCATCTAAGGGTAGACATCCCTTCAGTTGTGAAAAATCACAAATAAGGTCGATCGGATTTTAACTCA

154 E L K I L A D V G L V G F P S V G K S T L L S V I T S A K P X I G 186

801 CCTACCACTTTACCACTATTGTACCAATTTAGGTATGCTTCCACCAATCAGGTGAATCTTTGAGTAGCCGACTTGCAGGTTTGATTGAAGGGG 900
CGGATGTTGAAATGGTATACATGCTTTAAATCCATACCAAGCGTGGTTAGTCCACTTAGGAAAGCTCATCGGCTGAACGGTCCAACTAACTTCCCC

187 A Y H F T T I V P N L C M V R T O S G E S F A V A D L P G L I E G A 220

901 CTAGTCAAGGTGTTGGTTGGGAACTCAGTTCTCTCGTCACTCGAGCGTACACGTTTATCTTCACTCATTGATATGTCAGTACGGAAGGGCGTGA 1000
GATCAGTTCCACAACCAACCTTGAGTCAAGGAGGCGAGTGTAGTCCGATGTGACCAATAGGAAGTGTAGTAACTATACAGTCCGATCGCTTCCGCACT

221 S C G V G L G T C F L R N I E R T R V : L H I I D M S A S E G R D 253

1001 TCCATATGAGGATTACCTAGCTATCAATAAAGAGTGGAGTCTTACAATCTTCCGCTCATGGAGCGTCCACAGATTATTGTAATAAAGATGGACATG 1100
AGGTATACCTCCTAATGGATCGATAGTTATTTCTCGACCTCAGAAATGTTAGAGCGGAGTACCTCCGAGGTGTCTAATAACATTGATTATTCTACCTGTAC

254 P Y E D Y L A I N K E L E S Y N L R L M E R P Q : I V T N X M D M 286

1101 CTTGAGAGTCAGGAAAAATCTTGAAGAAATTAAGAAAAATTTGGCTGAAAAATATGATGAATTTGAAGAGTTACAGCTATCTTCCCAATTTCTGGATTGA 1200
GGACTCTCAGTCTTTTGAAGATTTCTAAATCTTTTAAACCGACTTTAATACTACTTAACTCTCAATGGTGGATAGAAGGGTTAAAGACCTAACT

287 P E S O E N L E E F K K F L A E N Y D E F E E L P A I F P I S G L T 320

1201 CCAAGCAAGGTCTGGCAACACTTTTAGATGCTACAGCTGAATTTGTTAGACAAGACACCAAGATTTTGGCTCTACGACGAGTCCGATATGGAAGAAAGT 1300
GGTTCTCTCAGACCGTTGTGAAATCTAGGATGTCGACTTAACAATCTGTTCTGTGGTCTTAAAAACGAGATGCTGCTCAGGCTATACCTTCTTCTCA

321 F O C L A T L L D A T A E L L D K T P E F L L Y D E S D M E E E V 353

Fig. 17 (Sheet 2 of 2)

1301 TTACTATGCAATTTGACGAAGAGAAAAACCTTTGAAATTAGTCGTGATGACGATGCGACATGGGTACTTTCTGGTGAAAAACTCATGAAACTCTTTAAAT 1400
AATGATACCTAAACTGCTTCTCTTTTGGGAAACTTTAATCAGCACTACTGCTACGCTGTACCCATGAAGACCACCTTTTGAGTACTTTGAGAAATTA
354 Y Y G F D E E E K A F E I S R D D D A T W V L S G E K L M K L F N 386

1401 ATGACCAACTTTGATCGTGATGAATCTGTCAATGAACTTTA 1441
TACTGGTTGAAACTAGCACTACTTAGACAGTACTTTGAAAT
387 M T N F D R D E S V M K L 399

gsp311

Fig. 18

(SEQ ID NO: 53) 1 TCGAATGCCCTTAAGAAAACAAATCGAAATCAAGAAAAACAGTAAGACAAGTTCTTTGTTCTATGAATTATTAGAAATGAAGAAAAGAGATATTAT 100
(SEQ ID NO: 54) 1 ACCTTAGCGGAAATCTTTTGTAACTTTAGTTCTTTTGTCTATCTGTTCAAGAAAAACAGTAATCTTAATAATCTTACTTCTTTCTCTATAATA 1
(SEQ ID NO: 52) 1 M 1

101 GGCTGAAGAAAGAGTAGAACCAAAACCAATTGACCTTGGTGAATATAAATTTGGTTTCCATGACGATGTAGAGCCTGTCTTATCGACAGGAAAGGACTC 200
CCGACTTCTTTCTCATCTTGGTTTGGTTAACTGGAAACCACTTATATTTAAACCAAGGTACTGCTACATCTCGGACAGAAATAGCTGTCTTTCTCTGAG 34

2 A E E R V E P K P I D L G E Y K F C F H D D V E P V L S T G K G L 34

201 AACGAAGGTGTTATTCTGTGAATTATCTGCTAAGGGTGAGCCTGAGTGGATGTTGGAGTTCCGTTTGAAGTCTTATGAAACCTTCAAAAAATGCCA 300
TTGCTTCCACAATAAGCACTTAATAGACGAGATTCCCACTCGGACTCACTCAAACTCAAGGCAAACTTCAGAATACTTTGGAAGTTTCTTACGGGT 68

35 N E C V I R E L S A A K G E P E W M L E P R L K S Y E T P K K M P M 68

301 TCGAAACTTGGGGAGCAGACTTGTGAGATTGACTTTGATGACTTAATCTACTACCAAAAAACCACTGCAAAACCAAGCCTGTTCTTGGGATGATGACC 400
ACGTTTGAACCCCTCGTCTGAACAGTCTCTAAGTGAAGTGAATAGATGATGTTTGGTAGACTGTTTGGTGGGGCAAGAACCTTACTACATGG 101

69 Q T W G A D L S E I D F D D L I Y Y O K P S D K P A R S W D D V P 101

401 TGAAGAAGATTAAAGAAACCTTTGAACGTATCGGGATTCCAGAAGCTGAACGTGCTTATTAGCAGGGGCTTCTGCCAGTACGAGTCAAGAGTGGTTTAC 500
ACTTTTCTAATTTCTTTGGAACTTGCATAGCCCTAAGGCTTTCGACTTGACCAATAAATCGTCCCGAAGACGGGTCACTCTCAGTCTTCAACAAATG 134

102 E K I K E T F E R I C I P E A E R A Y L A G A S A O Y E S E V V Y 134

501 CACAATCATGAGGAGAGCTTCCAAAAATAGGTATTATCTTACAGATACAGATTCCGCACTCAAGGAATACCCAGACTTATTAAACAATACTTTGGA 600
GTGTTGTAATCTCTCTCAAGGTTTTTAATCCATAATAGAAATGTCTATGTCTAAGGCGTGAGTTCTTATGGGTCTGAATAAATTTGTTATGAACCGCT 168

135 H N H K E E F O K L G I I F T D T D S A L K E Y P D L F K O Y F A X 168

601 AGTTGGTACCGCCGACAGATAACAAGTTGGCAGCCCTCAACTCAGCAGTATGCTCGGGTGGAACTTTTATCTACGTGCCAAAAGGTGTCAAGGTAGATAT 700
TCAACCATGGCGGCTGTCTATTGTTCAACCGTGGGAGTTGAGTGTCTATACCAAGCCCACTTGAAAAATAGATGCAGCGTTTCCACAGTCTCATCTATA 201

169 L V P P T D N K L A A L N S A V W S G G T F I Y V P K G V K V D I 201

701 TCCACTTCAAACTTATTTCCTATCAATAACGAAATATAGCTCAGTTCCAACTGATTATCGTTGATGAGGGAGCAAGCTCCACTACGTAGAA 800
AGGTGAAGTTTGAATAAAGCGATAGTTATTGCTTTTATATCCAGTCAAGCTTGCATGGAACTAATAGCAACTACTCCTCTGTCAGGTGATGCACTCT 234

202 P L Q T Y F R I N N E N : G O F E R T L I I V D E G A S V H Y V E 234

801 GGATGTACAGCACCAACATATTCAAGCAATAGCTTACAGCTGCCATTGTGAATAATTTTGGTTTGGACGGAGCTTATATGCGTTATACAACTATCCAAA 900
CTACATGTCTGTTGTATAAGTTCTTATCGAATGTGGCAGCGTAACATCTTTAAAAACGAAACCTGCTCGAATATACGCAATATGTTGATAGGTTT 268

235 G C T A P T Y S S N S L H A A I V E I F A L D G A Y M R Y T T I O N 268

901 ACTGGTCTGATAACGTCTATAACTTGGTAAACAAAGCGTGCTAAGGCTCAAAAGGATGCCACTGTTGAGTGGATTGATGGAACTTCCGTTGCCAAAACGAC 1000
TGACCAGACTATTGAGATATGAACCAATGTTTCCGACGATTCCGAGTTTCTCTACGGTGACAACTCACCTAACTACTTTCGAACCCACGGTTTCTCTG 301

269 W S D N V Y N L V T K R A K A Q K D A T V E W I D G N L G A K T T 301

1001 TATGAATATCCATCTGTTTACCTTGTATGGAGAGGAGCGGCTGATACCTGCTCTCTATCGCCTTTGCTAATGCGAGGCAACACCAAGACACGGGTGCT 1100
ATACCTTATAGGTAGACAAATGGAATACCTCTTCTCGCGCACCATGGTACGAGAGATAGCGGAAACGATTACGTCCGTTGCTGTTCTGTGCCACGA 334

302 M K Y P S V Y L D G E G A R G T M L S I A F A N A G O R O D T G A 334

1101 AAGATGATTCAAACTGCTCCACATACCACTGCTCTATTGCTCTAAATCATCGCTAAAGGTGGAGGAAAGGTTGACTACCGTGGACAACTCAGCTTTA 1200
TTCTACTAAGTGTACCGAGGTGATGCTCGAGCAGATAACACAGATTAGGTAGCGATTCCACCTCCTTTCCAACTGATGGCAGCTGTTCACTGGAAT 368

335 K M : H N A P H T S S S : V S K S : A K G G G K V D Y R G O V T F N 368

1201 ACAAGAACTTAAGAAATCTGTTTCCCACTTAATGTGATACCAATTATCATGATGACCTT 1263
TGTCTTCTGAGATTCTTACAGAAAGGTGTAACCTTACACTATGTAATAGTACCTACTGGAAA 1263

369 K N S K K S V S H : E C D T I I M D D L 388

qep3262 Fig. 19

(SEQ ID NO: 56) 1 AGCTGGGAATTTATGAGCAAGTATCCTATCTTAAGCAAGGAGAGTGTATCTAACTCGTTATAATCAAGTTCAAACTGAACAGCAACTTTAATCTTA 100
(SEQ ID NO: 57) TCGACCTTAAATACTCCTTCATAGGATAGAATTTCTTCTTTCACAAATAGATTGAGCAATATTACTTCAAGTTTGACTTTCTCGTTGAAATTAGAA
(SEQ ID NO: 55) 1 A G I Y E O V S Y L K E G R S V Y L T R Y N E V O T E T A T L I L 33

101 GGAGCTATTGTGGGATAGCTAGTTCCTTGTACTCTTTTATTCTGTCAATCTTCTATATTTCAGCAATTCGCGGAGATATCTTGATTAAACGAATTT 200
CCTCGATAACACCCCTATCGATCAAGGAACATGAGAAAAAAGACAGTTAGAAGATATAAAGCTCGTTAAGCGGCTCTATAGAACTAAATTCCTTAA
34 C A I V G I A S S L L L F Y S V N L L Y F E Q F R R D I L I K R I S 67

201 CAGGTTTACGATTTTGAACACATGCTCAGTATATGGTTAGTCAATTTGCCAGTTTGTATTTGGTGCTAGTCTCTTTATTTAAGCAGTCGAGACTT 300
GTCCAAATGCTAAAAAATTTGTACGAGTCATATACCAATCAGTTAAACGGTCAAAACATAAACCCAGATCAGAGAAATAAAATTCGTACGCTCTGAA
68 G L R F F E T H A O Y M V S O F A S F V F G A S L F I L S S R D L 100

301 GGTGATTGGCTTGCTCACTTTATTAGTCTTTCTAGCTAGTGCAGTTTTCAGCGCTTACCCTCAAGCGCAGAAAGAAATCTCGTCTTTCTATGACAAATTATG 400
CCACTAACCGAACCGAGTGAAATAATCAGAAAGATCGATCAGTCAAAACTGCGAAATGGCAGTTCGCGTCTTTCTTAGAGCACAAAGATACTGTTAATAC
101 V I G L L T L L V F L A S A V L T L Y R Q A Q K E S R V S M T I H 133

401 AAAGGAAAAATAGGATGATTGAACTAAAGAAATATATCTAAAAAATTTGGAAGCCGTCAGCTATTTTCAGATACGAATCTTTA 481
TTTCTTTTATCCTACTAAGTTCATTTCTTATATAGATTTTAAACCTTCGCGAGTCGATAAAAGTCTATGCTTAGAAAT
134 K C K * 117

gsp3387

Fig. 20

(SEQ ID NO: 59) 1 TTTTATCTAGTACAGTATATTTATGCGCTCTGCCCAATATTCAATCCATCCAAATGTATTAGAATGGATCTTAGTCTTCAAGATATGACGACTGG 100
AAAAATAGATCATGTTCATATAAATAACCGCAGCGGTTATAAGTTAGGTAGGTTTACATAATCTTACCTAGAAATCAAAATGAAGTCTTATCTACTGCTGACC
(SEQ ID NO: 60)
(SEQ ID NO: 58) 1 M T T G 4

101 AGTATATTGCTTTCCGTTCAATATATATTTCTTTTATTTGATGAATAACTATTTTAAAGGTTGGAGTGTCCGATTGCTGAAATCAATTAAG 200
TCATATAACGAAAGGCAAGTGTATATATAACAGAAAAAATAAATACTTATTGATAAAATATCCAACTCAGAGGTAAGCAGACTTTAGTTAATTC
5 V Y C P P F T Y I L F F F Y L M N W Y P N R L E C R I R L K S I K 37

201 CACTTACCAGTTTACTTTCAAATAGCAGCTCTTAGTACGGGATTGGACGGGCACTTTATTTTATTCATTTTCTAATTCATTAGTAATGGTT 300
GTGAAATGGTCAAAATCAAAGTTTAACTGTCGAGAATCATGCCCTAAACCTGCCCTGAAATAAAAATAAATAAAGATTAACTAAATCATTACCAA
38 H F T S F S F K L A A L S T C I W T A T L F L L I F L I A F S N G F 71

301 TTAGCTTCTCTTGGAGATAAAGGAGGTTGATTTTAAAGAGAAATTTATGGTATAAGTATTGCAACAAATGCTAGTTCTTTATAGGATTTTCTCTC 400
AATCGAAGAGAAACCTCTATTTCTCCTCAACTAAAAATTTCTTAAAAATACCATATTATAACGTTTGTACGATCAAGAGAAATATCCTAAAAAAGAG
72 S F S L E I K E V D F L R E F Y G I S I A W N A S F F I G F F F S 104

401 TTATATAGCATACTATTTCTTTTATCCTTACTTACTATTAGCAGTTTCTTGGTTAAAAAATCAACATGAGCTTAGTATTTCTGTTTACTTTTTTA 500
AATATATCGTATGATAAAGAAAAATAGGAATGAATGATAATCGTCAAAAAGAACCAATTTTATAGTTTGTACTCGAATCATAAAGACAAATGAAAAAAT
105 Y I A Y Y F F L S L L T I S S F S W F X K S N M S L V F L F T F L 137

501 TTTGTAGAATCCTTATCTGGATTTATCAGTTGGACAAATGGGATAATTGGATTATGCCAATTTTTCAGTATATGGTAAATTCCAATCCGTATGCATTGA 600
AAACATCTTAGGAATAGACCTAAATAGTCAACCTGTTACCTATTAACTAATAACGGTTAAAAAGTCATATACCATTTAAGGTTAGGCATACGTAAC
138 F V E S L F W I Y Q L D N G I I G L L P I F Q Y M V N S N P Y A L I 171

601 TTTATGGCTTACATTACTATCTATCATAATTCATTGACTGTAATTTCTGTTATAGAAAATGGAGGAGAGTGTAAAAGTTGGAAATGGGAAAGTTAAG 700
AAATAACCGAATGTAATGATAGATAGTATTAAGGTAACGACATAAAAGACAAGTATCTTGAECTCCTCTCAGATTTTCAACCTTTACCTTTCAATTC
172 Y W L T L L S I I I P L T V F S V H R N W R R V * 196

gcp4?

Fig. 21 (Sheet 1 of 2)

(SEQ ID NO: 62) 1 AGGGAACAGAAAAATTTGAGTTTTCGTGATATAATAGAGTCTGTATATAAGGAGGTAAATCATGGAGTTAGTGCAATGGAATTTCAAACATTTTATCC 100
(SEQ ID NO: 63) TCCCTGTGTTCTTTAAAGTCCAAAGCACTATATATCTTCAGACATATATCTCCCATTTAGTACCTCAATCACTGACCTTAAAGTTGTGAAAAATAGG
(SEQ ID NO: 61) 1 M E L V H G I S T H F I Q 11

101 AATCAAAAAAGTTTAAACAAACAAAATTACCGTGGTTTACCCTCCATTATCCCTGATACGATTGCAGGTACATGTTGAGTGCAAGTATGCTAGA 200
CTAGTTTTTCAAATTTGTTTGTTTTAAAGGCAAGCAAAATGGCGAGGTAATAGGGAATGCTAAGCTCAAGTGTACAACTCACTTCATACGATCT
14 S K R F K T N K I T V R F T A P L S L D T I A G H M L S A S M L E 46

201 GACTGCTAATCAGATGTACCCCACTTCTCAGATTTGAGGAGACACTTGGCCAGTCTATACGGTACAGATATGTCACCAATGTTTTCAGAAGAGGGCAA 300
CTGACGATTAGTCTACATGGGCTGAAGAGTTCTAACTCTCTGTGAACGGTCAAGATGCTATGCTATACAGTTGGTTAAACAAAGTCTTCTCCGTT
47 T A N Q M Y P T S O D L R R H L A S L Y G T D M S T N C F R R G Q 79

301 AGCCACATTATAGAAATGACATTTACCTATGTTCTGTGATGAGTTTAAAGTAGGAAAAACGCTAACCTCTCAGATTTTGGAACTGTGAAAAAGAACTC 400
TCGTTGTAAATCTTAACTGTAAATGGATACAGCACTACTCAAAAATTCATCCTTTTTCAGCAGATTGGAGAGTCTAAAACCTTGAACATTTTCTTTGAG
60 S H I I E L T F T Y V R D E F L S R K N V L T S O I L E L V K E T L 113

401 TTTTTCACCCGAGTGTGATAATGGGTTGATCCGGCTTATTTGAAATGAGAAAAACAAATGCTAGCAAGTTTAGCAGCTGATATGGATGATTC 500
AAAAAGTGGGCTCATCACTATTACCCAACTAGCCGGAATAAAGTTAACTCTTTTGTAAAGATGTTTCAAACTGTCGACTATACCTTAAG
114 F S P A V V D N G F D P A L F E I E K K O L L A S L A A D M D D S 146

501 TTTTATTTGCACATAAAGAAATGGATAAATGTTTTCATGATGAAGCTTCAATGGAAATAGTGAATTTACGAAATCGTATTTAGCTGAAACT 600
AAAAAATAAAGCTGTATTTCTTAACTATTAAACAAAAAGTACTACTCGAGAAGTTAACCTTATATCACTAAATGCTTTAGCATAAAAATCGACTTTGA
147 F Y F A K K E L D K L F F H D E R L O L E Y S D L R N R I L A E T 179

601 CCACAAAGTTCTTATTTCTGTTTCCAAGAAATTTAGCAATGATCGAAATAGATTTCTTTTCTAGGTGATTTTAAAGAGCTTGAATTTCAAATGTAT 700
GGTGTTCAGAAATAAGAACAAAGGTTCTAAAAATCGGTTACTAGCTTATCTAAAGAAAAAGGATCCACTAAAATTAAGCTTCAAGCTTTAAGTTTACATA
180 P Q S S Y S C F O E F L A N D R I D F F F L G D F N E V E I Q N V L 213

701 TAGAATCAATGGCTTTAAAGGTGCAAAAGCAGATGTGAAGGTTCAAGTATGTCACCTTATCTAATATCTTCAGGAAGGTATGGTTGGAAAAATGT 800
ATCTTAGTAACCGAAATTTCCAGCTTTCTCTACACTTCCAACTCATACAGTGGGAATAAGATTATAGGAAGTCTCTCCATACCAAGCCTTTTACCA
214 E S F G F K G R K G D V K V Q Y C C P Y S N I L Q E G H V R K N V 246

801 GGCACATCCATTTGGAATTAGGTTATCATACCTTCTAAATATGGTATGAGCAACATTTACCCATGATGTAATGAATGGTTTACTTGGTGGATTT 900
CCCTGTTAGGTAAAACTTAAATCCAATAGTAATGGCAAGATTATACCACTACTCGTTGTAATGGGTACTAACATTAAGTAACTTCAACCAATGAACCACTAAA
247 C O S I L E L G Y H Y R S K Y G D E O H L P M I V M N G L L G G F 279

901 GCTCACTTAAGCTCTTTACAAATGTCCTGAAATGCTGGATTAGCTTATACCAATTTCAAGTGAGCTTGATTTATTTAGTGGATTTCTTGAGGATGTATG 1000
CGAGTGAGATTCGAGAAATGTTTACAGGCACCTTTACGACCTAATCGAATATGGTAAAGTTCACTCGAATAAATAAATCACTAAGCACTCTTACATAC
280 A H S K L F T N V R E N A G L A Y T I S S E L D L F S G F L R M Y A 313

1001 CTGGTATCAATCGAGAAATCGTAACCAAGCTCGTAAATGATGAATCAACTGCTTGATTTAAAAAAGGTTATTTTACAGAGTTTGAGTTAAATCA 1100
GACCATAGTTAGCTCTTTAGCATTTGGTCCGAGCATTTTACTACTTATAGTTGACGAATAAATTTTCCAAATAAATGTCTCAAACTCAATTTAGT
314 G I N R E N R N O A R K M M N N O L L D L K K G Y F T E F E L N O 346

1101 CACCAAGGAAATGATTCCTTGGTGGTTGATTTCTCAAGATAATCAATCTTCATGATTGAAGCTGCTTATCAAAATGCTTATTTGAAAAATCTTCA 1200
CTGGTCTCTTACTAAGCAACCAAGCAATGAAGAGTTCTATTAGTTAGAGTAACCACTGCAAGAAATAGTTTACGGAATAAAGCTTTTGAAGT
347 T K E M I R M S L L L S O D N O S S L I E R A Y O N A L F G K S S 379

1201 GCAGACTTAAAGTTGCAATGCAAACTGAAACAAATGCAAAAGATGCTATTGTAGAGTAGCTAATAATGTGAAACTACAAGCCATTTACTTTATGG 1300
CGTCTGAAATTTCAACCTAACGTTTGAAGCTGTTAACTGTTCTACGATAAACATCTCATCGATTATTACACTTTGATGTTTGGTAAATGAATACC
380 A D F K S W I A K L E O : D K D A : C R V A N N V K L O A I Y F M E 413

Fig. 21 (Sheet 2 of 2)

1301 AAGGAATAGAATGACAAAGGTTGT...TTGAAGAAAAATACTATCCAGCTGTAAAGAAAAAGGTTTATCGAACTCGTTTGGCCAAACGATTGACAGTTGCT 1400
TTCTTATCTTACTGTTTCCAAACAAAACTTCT...TTTATGATAGGTCGACATTTTCTTTTCCAAATAGCTTGAGCAAAACCGGTTGCCTAACTGTCAACGA
414 G I E * 417

gcp61

Fig. 22

(SEQ ID NO: 65): GTTTTGGACCATTTCAAAGTCGTTAGCAGAGAAAGAGTCGTTATCTTCCGAAAGAAATTTATACCTTTCAAAATCTGACTTTGGTATTTATTT
(SEQ ID NO: 66): CAAAAAAGCTGTAAGCTTTTCAGCAATCGTGCTTTTCTTCAGCAGATATGAAGCTTTCTTTAAATAATGGAAGTTCTAGACTGAAACCATAAATAAA 100

101 TAGAGAAAAATTAAGTTCTCCCATGCTTTATGAGAGGTTCTCTGTTATGCCAAATGAAGATTTAGTAGTGGAAATCTGGGAAATGACTCCCAAAACAAAGT 200
ATCTCTTTTAAATTCAGAGGGTACCAAAATACCTCTCCAGGACAAATACGCTTACTTCTAAATCATCACCTTAGACCTTTAACTGAGGGTTTGTTCAT

(SEQ ID NO: 64): M V Y G E V P V Y A H E D L V V E S G K L T P K T S 26

201 TTTCAATAACCGAGTGGGCTTAAATAAACAGGAATTCAGTATTTAAGCTATCAAAATCATCAATTTATAGCTGGGACAAACGATTTTATATGATC 300
AAGTTTATGGCTCACCGCAATTTATTTGCTTAAAGTCATAAATTCGATAGTTTAGTAGTTAAATATCGACGCTGTTTGTCAAAAAATATACTAG

27 F Q I T E W R L N K O G I P V F K L S H Q F I A A D K R F L Y D Q 60

301 AATCAGAGCTAATCCCAAAATAAAAAAGTATGGTTAGAAATCTGACTTTAAACTGTACAAATAGTCTTATGATTTAAAGAAAGTGAATCATCTTTATC 400
TAGTCTCCATTTAGGTTGTTATTTTTCATACCAATCTTAGACTGAAATTTGACATGTTATCAGGAATCTAAATTTCTTCACTTTAGTAGGAATAG

61 S E V T P T I K K V W L E S D F K L Y N S P Y D L K E V K S S L S 93

401 AGCTTATTCCCAAGTATCAATCCACAGACCATGTTTGTAGAAGGAAGAGAAATTTCTACATATTGATCAGGCTGGATGGGTAGTAAAGAAATCAACTCT 500
TCGAATAAGCGTTATAGTTAGCTGTTCTGGTACAAACATCTTCTCTTAAAGATGTATAACTAGTCCGACCTACCCATCGATTTCTTAGTTGAAGA

94 A Y S O V S I D K T M F V E G R E F L H I D O A G W V A K E S T S 126

501 CAAGAAGATAATCGATGAGTAAAGTTCAAGAAATGTTATCTGAAAAATATCAGAAAGATTTCTTCTATTATGTTAAGCACTGACTACTGAAAAAG 600
CTTCTTCTATTAGCTACTCATTTCAAGTTCTTCAATAGACTTTTATAGTCTTTCTAAGAAAGAGATAAAATACAAATTCGTTGACTGAGCTTTTC

127 E E D N R M S K V Q E M L S E K Y O K D S F S I Y V K Q L T T G K E 160

601 AAGCTGCTATCAATCAAGATGAAAGATGATGCGCCAGCGTTTTCGAACTCTCTTATCTCTATTATACGCAAGAAAAATAAATGAGGCTCTTTATCA 700
TTGACCATAGTTAGTTCTACTTTCTACATACCTCGGTCCGAAACTTTGAGAGAAATAGAGATAATATGCTTCTTTTATTACTCCGAAATAGT

161 A G I N O D E K M Y A A S V L K L S Y L Y T O E K I N E G L Y O 193

701 GTTAGATACGACTGTAAATACGTTATTCAGTCAATGATTTTCAGGTTCTTATAAACAGAGGGAAGTGGTAGTCTTCTAAAAAGAGATAATAAA 800
CAATCTATGCTGACATTTATGATAGACGTCAGTTACTAAAGGTCGCAAGAAATTTGGTCTCCCTTACCATCAGAAAGGATTTTCTTCTATTATTT

194 L C T T V X Y V S A V N D F P G S Y K P E G S G S L P K K E D N K 226

801 GAATATTCTTAAAGGATTTAATTACGAAAGTATCAAAAGATCTGATAATGTAGCTCATATCTATTGGATATTACATTTCAACCAATCTGATGCCA 900
TCTATAAGAAATTTCTAAATTAATGCTTTCTATAGTTTCTTAGACTATTACATCGAGTATTAGATAACCTTAAATGTAAGTTTGGTTAGACTACGGT

227 E Y S L K D L I T K V S K E S D N V A N N L L G Y Y I S N O S D A T 260

901 CATTCAAATCCAGATGCTGCCCATTATGCGAGATGATGGGATCCAAAGAAAAATGATTTCTTCTAAGATGCGCGGGAAGTTTATGGAAGCTATTTA 1000
GTAAGTTTAGGTTCTACAGACGGTAATACCTCTACTAACCTAGGTTTCTTTTAACTAAGAAAGATTTACCGGCTTCAATACCTTCGATAAAT

262 F K S K M S A I M G D D W D P K E K L I S S K M A G K F M E A I Y 293

1001 TAATCAAAATGGATTTGTCTAGAGTCTTTCAGTAAACAGATTTTGTAGATCAGCGAATTCGCAAGGTTCTTCTTAAAGTAGCTCATAAATTTGGA 1100
ATTAGTTTCTTAAACAGGATCTCAGAACTGATTTGTCTAAACTATCAGTCTGCTTAACGGTTTCCCAAGAGACAAATTCATCGAGTATTTAACT

294 N C N G F V L E S L T K T D F D S Q R I A K G V S V K V A H K I G 326

1101 GATGCGGATGAATTAAGCATGATACGGGTTGTCTATGCAGATTCCTATTTATTCTTTCTATTTTCACTAAGAAATCTGATTATGATACGATTTCTA 1200
CTACGCTACTTAAATCTGACTATGCCCAACAGATACGTTAAGAGGTAAATAAGAAAGATAAAGTGATTTCTAAGACTAACTATGCTAAGAT

327 C A D E F K H D T G V V Y A D S P F I L S I F T K N S D Y D T I S K 360

1201 AGATAGCCCAAGGATTTTATGAGGTTCTAAATGAGGGAACCAATTTTAAATCAATTTCTCAAGAAAGGATATTTCAAAAGCATGCTAAGGCGGT 1300
TCTATCGGTTCTTCAAAATCTCCAAGATTTACTCCCTTGGTCTAAAAAATTTAGTAAAGAGTTCTTCCCTATAAGTTTTCGTAGGATTCGCCCA

362 : A K D V Y E V L F * 371

gap76

Fig. 23

(SEQ ID NO: 68) 1 TTGAAAAATATTATCTATAAGAACGACATATAAATGTAA CAAAGCGTAAATATTATTAGGCTTTTTCGTATACTAGTATTGCTTTTAAAGAAAGGA 100
(SEQ ID NO: 69) AACTTTTATAATAGATATTCTTGCTGTATATTACATTGTTTCCGCATTATAAATAATCCGAAAAAACCATATGATCATAACAGAAATTTTCTTCCT

101 GTATCTAGCTAATATGAAGAAAAAATCTTAGCGTCACCTTTTATTAAGTACAGTAATGCTTTCTCAAGTAGCTGTTTAAACAACTGCCATGCGAGAAACG 200
CATAGATGCAATTATCTCTTTTATAGAAATCGCAGTGAAAAATATTATGTCATTACCAAGAGTTTATCGACAAAAATGTTTACCGCTACGCTCTTTCG

(SEQ ID NO: 67) 1 M K K K I L A S L L L S T V M V S Q V A V L T T A H A E T 29

201 ACTGATGACAAAAATGCTGCTCAAGATAATAAATTAGTAACCTTAACAGCACAAACAAGAGCCCAAAAAACAAGTTGACCAAAATTCAGGAGCAAGTAT 300
TGACTACTGTTTAAACGACGAGTTCTATTATTTTAAATCATTTGAATGTCGTGTTGTTGTTCTTCGGGTTTTTGTTCAACTGGTTTAAAGTCTCTGTTGATA

30 T D D K I A A Q D N K I S N L T A Q Q Q E A Q K Q V D Q I Q E Q V S 63

301 CAGCTATTCAAGCTGAGCAGTCTAACTTCCAAAGCTGAAAAATGATAGATTACAAGCAGAACTTAAGAAACTCGAGGGTGAGATTACAGAACTTTCTAAAAA 400
GTCGATAAGTTCCACTCGTCAGATTGAACGTTTCACTTTTACTATCTAATGTTGCTCTTAGATTCTTTGAGCTCCCACTCTAATGTCTTGAAGATTTTT

64 A I Q A E Q S N L O A E N D R L O A E S K K L E G E I T E L S K N 96

401 CATTTGTTCTCGTAACCAATCGTTGAAAAACAAGCTCGTAGTGCTCAAAACAATGGAGCCGTAACCTAGCTATATCAATACCAATGTAACCTCAAAATCA 500
GTAACAAGAGCATTGCTTAGCAACCTTTTGTTCGAGCATCAGGAGTTTGTTCCTCGGATTGATCGATATAGTTATGGTAACATTGAGTTTGTAGT

97 I V S R N Q S L E K O A R S A Q T N G A V T S Y I N T I V N S K S 129

501 ATTACAGAAGCTATTTCACTGTTGCTGCAATGAGTGAATCGTATCTGAAACAACAAATGTTAGAAACAACAAAGCCAGATAAAAAGCTATTTCTG 600
TAATGTTCTTCGATAAAGTGCAACGACGTTTACTCACTTTAGCATAGAGCTTTGTTGTTTACAACTCTTGTGTTTCCGCTATTTTTCGATAAAGAC

130 I T E A I S R V A A M S E I V S A N N K H L E Q Q K A D K K A I S E 163

601 AAAAAAAGTAGCAAAATAATGATGCTATCAATCTGTAATTGCTAATCAACAAAAATGCTGATGATGCTCAAGCATTGACTACGAAACAGGCAGAACT 700
TTTTTGTTCATCGTTTATTACTACGATAGTTATGACATTAAAGATTAGTTGTTTTTAAACCGACTACTACGAGTTTCGTAACGATGCTTTGTCGGTCTTGA

164 K O V A N N D A I N T V I A N O O K L A D D A Q A L T T K O A E L 196

701 AAAAGCTGCTGAATTAAGTCTTGTGCTGAGAAAGCGACTAGCTGAAGGGGAAAAAGCAAGCTATTAGAGCAAGAGCAGCTCAGGCAGAGGCTCG 800
TTTTCCAGCACTTAATTCAGAACGACGACTTTTGGCTGATCGACTTCCCTTTTTTGGTTCCGATAATCTCGTTCTTCGCTCGCTCCCTCTCCGAGC

197 K A A E L S L A A E K A T S 211

Fig. 24

YNES_BACSV

(SEQ ID NO: 71) 1 ATGTTAATGCTTTATTGATTATTTGGCTACTTGTATAGGCAGCATTCCATCTGGCTTAATTGTGGCAAGCTTCGCAAGGAATTGATATTGGGAGC 100
(SEQ ID NO: 72) TACATTAACGAATAACTAATAAAACCGATGAACATATCGGTGTAAGGTAGACCGAATTAACACCGCTTCGAACGGTTTCCTTAACATAAGCCCTCG
(SEQ ID NO: 70) 1 M L I A L L I I L A Y L I G S I P S G L I V G K L A K G I D I R E H 34

101 ACGGAAGCGGCAACTTAGGCGCTACCAATGCATTCCGTACATTOGGTGTAAAGCTGGTTCGGTCGTATAGCCGAGATATTTGAAAGGCACACTGGC 200
TGCCTTCGCGTTGAATCCGCGATGGTTACGTAAAGCATGTAAACCACATTTTCGACCAAGCCAGCAGTATCGGCCTCTATAAAAATTTCCCTGTGACCG
35 G S G N L G A T N A F R T L G V K A G S V V I A G D I L K O T L A 67

201 AACTGCATTCGCTTTTCTCATGCGATGTGATATTCACCGCTTCTTGCAGGAGTCTTTGCGGTTTTAGGCCACGTGTTTCCCATCTTCGCCAAATTTAAA 300
TTGACGTAAACGAAAAGAGTACGTACAACTATAAGTGGCGAAGAACGTCTCAGAAACGCCAAATCCGGTGCAAAAGGAGTAGAGCGGTTTAAATTT
68 T A L P F L M H V D I R P L L A G V F A V L G H V F P I F A K F K 100

301 GCGGTAAAGCCGTGCGGACATCAGGAGCGTTTTGCTATTTACGCACCCCTGTTATTTATCAGGATGGTTGCGGTATTTCTCATCTTTTATACTTGA 400
CCGCCATTTTCGGCACCCTGTAGTCTCTCCGCAAAACGATAAAATGGTGGGCAATAAATAGTCTACCAACGCCATAAGAAGTAGAAAAATATGAACT
101 G G K A V A T S G G V L L F Y A P L L F I T H V A V F F I F L Y L T 134

401 CTAATTTGTTTCTCTCATGATGTTAAGCGGATCTATAGTCTTATATATAGTCTTCTTGTCCATGATACGTATTTATTGATTGTGTTACCCCTGCT 500
GATTTAAACAAGAGAGAGTAGCTACAATTTGTCCTAGATATGACAAATATATATCAAGAAACAGGTACTATGCATAAATAACTAACAGCAATGGGACGA
135 K F V S L S S M L T G I Y T V I Y S F F V H D T Y L L I V V T L L 167

501 CACTATTTTGTGATATACAGACACCGAGCGAACATTAAACGAATTATCAATAAAACAGAACCTAAAGTAAATGGTTATAA 582
GTGATAAAAACACTATATGCTGTGGCTGCTGTAAATTTGCTTAATAGTTATTTTGTCTTGATTTTCATTTTACCAATATT
168 T I F V I Y R H R A H I K R I I N K T E P K V K W L * 193

Strategy for the targeted deletions of genes in *S. pneumoniae*

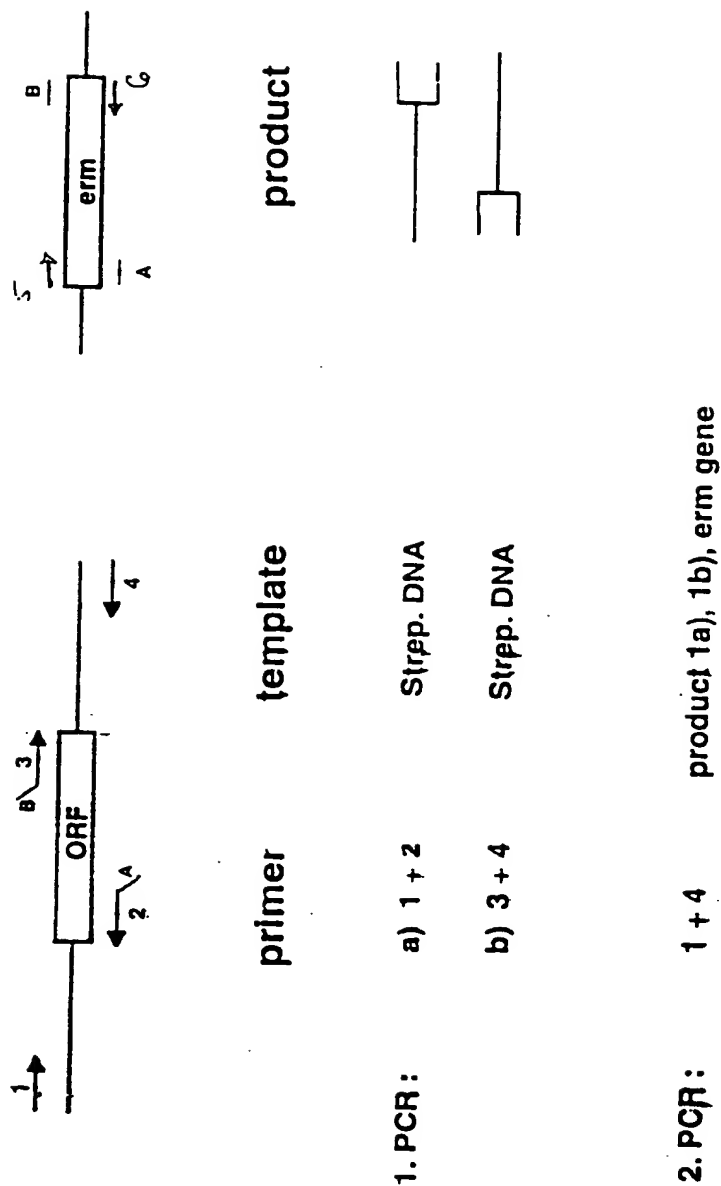


FIG. 25

Non-polar gene knockouts in *S. pneumoniae*

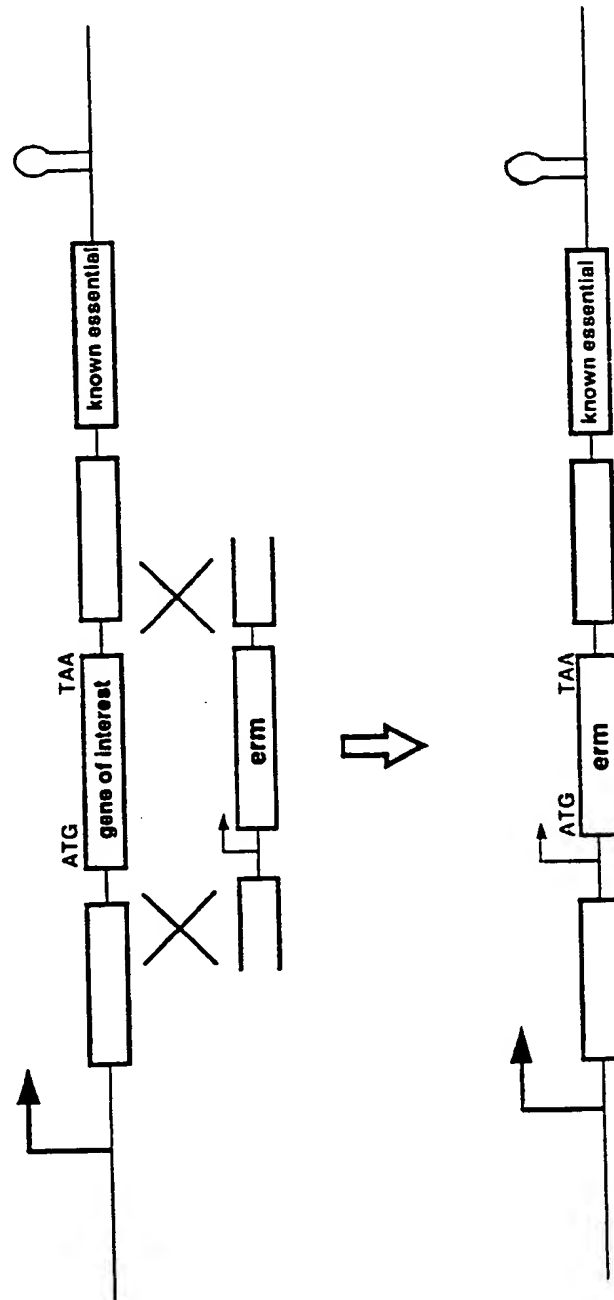


FIG. 26

- 1 -
Sequence Listing

gcp101

(SEQ ID NO: 2) 1 TGCTGATTTTGGAGAAAGTTTATTAGAGATAAAAGAGTCTAAGGAAAAAATTCATTTCATATTTTCTTCTATAAAATAGATAAAATCGTACAAAT 100
(SEQ ID NO: 3) ACGCTAAAAACCTCTTCAAATAATCTCTATTTCTCAGATTCTCTTTTAAAGTAACTATAAAAGAAGATATTTATCTATTTTACCATGTTAT

101 ATAAATTGAGGTAAATAAGGATGAGATTAGATAAAATATTTAAAGTATCGCGAATTATCAAGCGTCGTACAGTCGCAAGGAAGTAGCAGATAAAGGTAGA 200
TATTTAACTCCATTATTCCTACTCTAATCTATTTATAAATTTTCATAGCGCTTAATAGTTCCGAGCATGTCAGCGTTTCCTTCATCGTCTATTTCCATCT

(SEQ ID NO: 1) 1 M R L D K Y L K V S R I I K R R T V A K E V A D K G R 27

201 ATCAAGCTTAATGGAAATCTTGGCCAAAAGTTCAACGGACTTGAAAGTTAATGACCAAGTTGAAATTCGCTTTGGCAATAAGTTGCTGCTTGTAAGTAC 300
TAGTTCCAAATACCTTAGAACCGGTTTCAAGTTGCGTGAACCTTCAATTACTGGTTCACTTTAAGCGAAACCGTTATTCACGACGAACATTTTCATG

28 I K V N G I L A K S S T D L K V N D Q V E I R F G N K L L L V K V L 61

101 TAGAGATGAAAGATAGTACAAAAAAGAGATGCAGCAGGAATGTATGAAATTATCAGTGAAACACGGGTAGAAGAAAAATGCTAAAAATATTGTACAA 400
ATCTCTACTTTCTATCATGTTTTTCTTCTACGTCGTCCTTACATACCTTAATAGTCACTTTGTCGCCATCTTCTTTACAGATTTTATAACATGTTA

62 E M K D S T K K E D A A G M Y E I I S E T R V E E N V * 89

gsep1119

(SEQ ID NO: 5) : GAAATCTCTTTCGAATGTGACTGTAGCCATGAACGCTTATCAACGCTTTGCGACGCTTCCAGCTCAGACTTACAGGAAATGAAGAGCAAGACCACTG 100
(SEQ ID NO: 6) : CTTTAGGCAAGGTTACCTGACATCGGTACTTGGAAATATCTTGGAAACGCTCGGAAGGTTCCAGTCTGAATGCTCTTACTTTCTCTCTGCTGCTG

101 GGGCAGAAATCACTTGTCAATTTGTCGCAAACTACTTACAACCTTTGATGAAAGGACCTGGAGGAACTCATTGCTGACAAATCTTAATACACCTTTATGA 200
CCTGCTCTTAACTGAACAGTTAAGACGCTTTCATGAATGTTCAAACTACTTTCTCTGACCTCTTTCAGTAAGCACTGTTTAGAATTTATGTCGAAATACT

(SEQ ID NO: 4) : M K R T W R N S F V T H L N T P P H I 19

201 TTGGCAATATTGAGATTCCCAATCTGACCTTTTACCGCTATGGTGGGCTGACCAACTCAGCTTTCTGACCATGCGAAAGAGCTCGGAGCTGCACT 300
AACCTTTATAACTTAAGGCTTAGCATGCGAAATCGCGGATACCGACGCACTGTTGAGTGGGAAAGCATGGTAGCTTTTCTGAGCTCGAACTGA

20 G H I E I P N R T V L A P M A G V T H S A F R T I A K E L G A G L 52

301 CTTGTAAATGGAAATGCTCTGCAAGGCAATCCAAATACAAACGAAAGAAACCTGCAATGCTTTCATATGATGAGGCGGAAAGCCCTGCTCTATC 400
GCAACATTACTTTACGAGAGCTGTTCTCTTAGGTTATGTTGTTGCTTTTTTGGAGCTATACGAAGTATAGCTACTCCCGCTTTTGGGACAGAGATAG

53 V V M E M V S D K G : Q Y N N E X T L H M L H I D E G E N P V S : 85

401 CAATCTTTGCTAGCGATGAAGACAGCTTAGCAACGCGCAGGAGAAATCATCAAGAAACACCAAGACCGATATCGTGGATATCAACATGGGCTGCTCTG 500
GTTGAAAGACCATCGCTACTTTCTGCGATCGTGGCTGCTCTTAAGTAGCTTTCTTGTGCTTCTGGCTATAGCAGCTATAGTTGTACCCGACGGGAC

86 C L F G S D E S L A R A A E F : G E N T X T D I V D I N H G C P V 119

501 TCAACAAATCTGTAAGAACGAAGCTGGAGCTATGTTGCTCAAGCATCTGCAAGATCTACTTATCATCAACAAAGCTCCAGTCTGCTCTGATATCCC 600
AGTTGTTTACGACTTTCTGCTTGCACCTGATACACCGAGTCTTAGGACTGTTCTAGATGAGATAGTAGTTGTTTCCAGTCCAGACGGAAGCTATAGGG

120 N K : V K M E A C A M M L K D P D K I Y S I I N K V Q S V L D I P 152

601 ACTTACTGTCAAAATGCTTACCGCTTGGCGGACCATCTCTGGCAGTAGAAATGCTCTGCTGAGCTGCAAGTGTCTGCTGCTTCCGCTCGCCATGCA 700
TGAATGACACTTTACCGCATGGCGGACCTCGCTGGTAGAGACCGTCACTTTTACGGGAGCGACGACTCGGACGTTCCAAAGACGGGAGCGGTACGTA

153 L T V K M R T G W A D P S L A V E N A L A E A A G V S A L A M H 185

701 GGGCTTACCGGTGAACAAATGTATCTGGCCACCGACCTTGACACTTTTCAAGGTTGCCCAGCTCTAACCAAGATTCCATTTCATCGGCAACGCTG 800
CCGGCATGGGCACTTGTATACATATGACCGGTGCTCTGGAACTCTGGGAAATGTTCCAAACGGTTCCGAGATTGTTCTAAGGTAAGTAGCGGTTGCCAC

186 G R T R E O M Y T G H A D L E T L Y K V A O A L T K I P F I A N G D 219

801 ATATCGCTATCTGCAAGAGCCAGCAACCATCGAAGAAGTTGCTGCTGACCGCTCATGATTGGCCGAGCTGCCATGGGAAATCTTACCTCTTCAA 900
TATAGGCTAGACAGCTTCTCGCTTCTGCTAGCTTCTTCAACCAACGACTGCTGAGTACTAACCGGCTCGACGGTACCTTTAGGAATGGAGAGTT

220 : R T V Q E A K C R I E E V G A D A V M I G R A A N G M P Y L F N 252

901 CCAATCAACCATTACTTTCAAAACAGGAGAAATCTTACCTGATTTGACCTTTGAAGACAAGATGAAGATCGGCTACGAACACTTGAACAGATTGATTAAC 1000
CGTTTAACTGGTAATGAACCTTTCTCTTTTAGGATGGACTAACTGGAACCTTCTGTTCTACTTCTAGCGGATGCTTGTGAACCTTTGCTAACTAATTG

253 Q I M H Y F E T G E I L P D L T F E D K M K I A Y E H L K R L I N 285

1001 CTCAAGAGGAGAAACCTCGGAGTTCGTGAATTCGCGGCTCTGCTCTCACTATCTCCGTGCAACATCTGGCGCTGCCAACTCCGTGGAGCAATTTCCG 1100
GAGTTTCTCTTTTGCAGGCTCAAGCACTTAAGGCGCGGAGCGAGGAGTGTATAGAGGCACTTTGTAGACGGGAGGTTTGGAGGCACTTCGTTAAAGCG

286 L K G E N V A V R E F R G L A P H Y L R C T S G A A K L R O A I S O 319

1101 AAGCTAGCACCTTAGCAGAGATTGAAGCCCTTTCGAATTCGAGAAGGCTTAATAGTTTAAACCCGTAACCTCTCTTAAGAGTCTCTGAATGCCGCA 1200
TTCCATCGTGGCATGCTCTTAACCTTGGGAGAACCTTAACCTCTTCCGAATTAATCAAAATTTGGCATTGAGAGAAATTTCTCAGAGAACTTACGGCGGT

320 A S T L A E I E A L L O L E K A * 336

gcp1122

(SEQ ID NO: 8) : AAGGCAACGAGCTGGAAGTTCCTCTCATATTTTCAATAGTTATTAGCTACAGTTGAGCAACTTCAGAAAAATCAAAATTCCTCTCAAGTTCCTCTCTA 100
 (SEQ ID NO: 9) : TCCGTGCTCGACCTTCAAAAGGGAGTATAAAAAAGTTATCAAAATAATCGATGTGCAACTCGTTGAAGTCTCTTAAAGAAAGTTCAGAGAGAAGAT

101: TAGTAGATTTCGAAATCCCTTTTCAGCTAGTTTCTGASTCAGCACATAAGGACCTTTCTCTCTGAAAGTTGATTGGTATTGATGATAGCATAAAGCTTA 200
 ATCATCTAAAACTTTAGGAAAAAACTCGATCAAGAGCTCAGTCTGTATTTCTGGGAAACAGAGGACTTTCAACTAACCAATAACTACTATCGTATTGGCAT

201: CTGACCATCATTAATCCACTTATCTCTTAAGATTAGCAATAACTTCAGAAACGATGTTTTATCAATATGTTATTTTTCAGATATTCTCTGACTTCT 300
 GACTGCTAGTAATTAGGTGAATAGAAGAAATCTAATCGTTATTGAAGTCTTCTCTACAAAAATAGTTATAGCATAAAAAGTCTATAGAGACTGAAGA

301: TTTTCAGTGCCTGCTTTAAAGCATAAAGTGTAGAGGGCCAGATTCTTACCATAAGAAAAATGAGCAAGTCTTGAATCTCTTCAATTCCTCTTGGCTTA 400
 AAAAGTCAGGCAGGAAATTTCTATTCACCATCTCCGGTCTAAGAATGTTATTCTTTAACTCGTTTTCAGAACTTAGAGAAAGTTAAGGAGAGCGAAT

401: TCACCTTATCTCTCGATAACAATAAAACGAACAATTTGTATCTTCCGTGATATAGCATTTGTGCCCAATTATCAAGCTCCATCAGATAGAGTCTTTTCTT 500
 AGTGAATAGAGAGCTATTGTATTTCTCTTAAACATAGAAGCCATATATCGTAAAACAGCGGTAAATAGTTCCAGGTAGTCTATCTCAGAAAAAAGAA

501: TCAAGTTCCTGATTTTCATAGCTCTATTATAACTCAAAATGTGATAGATAGCGGTATGAATCTCAAGTCAAAACAAAAATACCAATAAAAATCAAG 600
 AAGTCAAAACACTAAAAGTATCGAGATAATATTCAGTTTACACTATTCTATCCCATACTTAGACTTTCACTTTGTTTTATGGTAAATTTTATGTTTC

(SEQ ID NO: 7) : M N L K V K Q R I P L K I K 14

601: CCGATCGGAATTAACCGTGAGCGAATCCGCTTTACAAAAAACATTACTTTTGTACAGGAGCTCTCAAGCGGAGATATCTATTGTCAGATTACTT 700
 CGGTACCTTAATTTGCACTCTCTTAGCGGAAATGCTTTTGTAAATCAGAAACATGCTCTCGAGAGTTTCCGCTTCTATAGATTAACAGTCTAATGAA

15: R H G : N G E C : G F Y Q K T L V F V P G A L K G E D I Y C O I T S 48

701: CTATTAGACGCACTTTGTCAGCAAAAATTAAGTCAAGGTCAACAAGAGCTCAAAATTCGAATTTGTCATCTCTCTACTATTATTAATGAATCGCGAGC 800
 GATAATCTCGCTTGAACAACTTCTTTAATGACTTCAAGTTGTTTTCAGATTAAAGGTTAACACGGTAGAACATGATAAATATTAATTAAGCTTCT

49: : R R N F V E A K L L K V N K K S K F R I V P S C T I Y N E C G G 81

801: CTGCAAAATCATGCACTGCAATTAAGTAAAGCAGCTGAGTTCAGACCGGACTTACTTCAACCGGCGAAAAAATTTGCTCTGCGAGGATATGAATA 900
 GACGGTTAGTAGCTGAGCAATATCTATCTGCACTCAAGTTCTGCTGAATGAAGTAGTTCGGGACTTTTAAAGCAGGACGCTCTATAGTTTTTA

82: C G I M H L H Y D K C L E F K T D L L N O A L K K F A P A G Y E N 114

901: TATGAAATTCCTCAACTATTGGAATGCAGCAACCAAAATATTACAGAGCTAAGTTACAAATTCAGACTCGAAAAATTAATAATCAAGTCAAGCGCGCT 1000
 ATACTTTAAGCAGGTTGATAACCTTACGTCTTGTATTATAATGCTCTGATTCAATGTTAAAGTCTGAGCTTTTAAATTTTATGTCAGTTCTCGCGCGA

115: Y E I R P T I G M O E P K Y Y R A K L O F Q T R K F K N O V K A G L 148

1001: TATATCCAAAACTCTCACTATTAGTAGAGTTGAAGACTGCTGTGATCAAGATAAGCAAAACCAAGTGAATGCTAAATCCCTTAGCAGAAATTAATAC 1100
 ATATACCTGTTTTGAGAGTGATAAATCATCTCAACTTTCTGAGGACCATGTTCTATTCTTTGGGTTCACTAACGATTAGCGAATCOTCTTAATGAATG

149: Y A O N S N Y L V E L K D C L V O D K E T Q V I A N R L A E L L T 181

1101: TTATCACCAGATTCCAATCAGCGATGAGAGAAAGTTCTAGGTGCTGCTACTATTATGCTCGAGCGCGGAGAAAGACCGGACAGGTTCAAGATTATTAT 1200
 AATAGTGTCTAAGGTTAGTGCCTACTCTTTTCAAGATCCACAGGCAATGAATAACCAAGCTCGCGGCTCTTTCTGCGCTTCCAGTCTAATAATAA

182: Y N G I P I T D E R K V L G V R T I M V R R A R K T G O V O I I I 214

1201: GTTACAAACCGCCAGCTTAATTTAACTCAATTCGTAAGAGAGTTGTTAAAGATTTCCAGAAAGTTGTGACAGTAGCTGTTAATACAAATACAGCTAAAA 1300
 CAATGTTTGGCGGTGCAATTAATTAAGTTAACCATTCTCAACCAATTTCTAAGGGGCTTCAACACTGTCTGACAAATTAATGTTATGTCGATTTT

215: V T M R O L N L T O L V K E L V E D F P E V V T V A V N T N T A R T 248

1301: CCACTGAGATATATGCTGAAAGACAGACATTATCTGGGGCAAGAGAGTATCAAGAGGTGTTACTCAATTAATGAATTTTCACTATCCCTCGAGCTTT 1400
 GGTCACTCTATACCACTTTCTGCTCTAATAGACCCCGTTCTCTCAATAGTTCTTCCACATGAGTTAATACTTAAAGTGAATAGGAGAGCTGAAA

249 S E I Y G E K T E : I M G Q E S I O E G V L N Y E F S L S P R A F 281

1401 TTATCACTAAATCCTGAGCAACAGAACTCCTCTATACCGAAGCAGTAAAGCGCTGGATGTTGATAAAGAACACCATTTGATTGACGCTTATTGTGCA 1500
AATAGTTGATTTAGGACTCCTTTGTCTTCAGCAGATATGGCTTCGTCTTTTCGGACCTACAACTATTTCTTCTGGTAAACTAACTCGGAATAACACCT

282 Y O L N P E O T E V L Y S E A V K A L D V D K E D H L I D A Y C G 314

1501 GTTGGACCATTTGCAATTTGCTTTTTCAGAAAGTAAAACTCAGACCTATCGATATTATTCAGAAAGCTATTGAAGATGCCAAGCGAAATGCTAAAA 1600
CAACCTTGCTAACTAAACCGAAACGTTTCTTCACTTTGTGAGTCTCCATACCTATAAAGGTCTTCGATAACTTCTACGGTTCTGCTTACGATTTT

315 V G T I G F A F A K K V K T L R G M D I I P E A I E D A K R N A K R 348

1601 GAATCGCATTTGACAACTACTCATTATGAAGCTGGAAACCGCAGAAAGATTATTCTCTGTTGTTACAAGGAAGCTACCGACAGCATGCTTTGATTGTTGA 1700
CTTACCTTAAACTGTTATGAGTAATACTTCGACCTTGGCTCTTCTAATAAGGAGCAACCATGTTCTCTCCGATGGCTCTGCTACGAACTAACCACT

349 M G F D N T H Y E A C T A E E I I P R W Y K E G Y R A D A L I V D 381

1701 CCCACCACTACAGGTCTGGATGATAAGTTATTAGATACTATTCTTACTTATGTACCAGAAAAATGGTTTATATTCTTGTAAATGTTTCCACCTTGGCT 1800
GGGTGGTGCATGTCAGACCTACTATTCAATAATCTATGATAAGAAATGAATACATGGCTTTTTTACCAGATATAAAGAACATTACAAAGCTGGAACCGA

382 P P R T G L D D K L L D T : L T Y V P E K M V Y : S C N V S T L A 414

1801 CCGTATTTGGTACGCTTAGTAGAAGTCTATGATCTTCATTATATCCAGTCGGTCCATATGTTCCACATACAGCTCGAAGCTGAAGCTGTTGTAATAATTAA 1900
GCACTAAACCATGCGAATCATCTTCAGATCTAGAAGTAATATAGGTCCAGCCAGCTATACAAGGGTGTATGTCGAGCTTGACTTCGACAACTTTTAATT

415 R D L V R L V E V Y : L H Y I O S V D M F P H T A R T E A V V K L I 448

1901 TAACAAAAGTTTAAAAAGTACTTCACAAAAGTTTGAAGAGCTGTATATAGTAAGAGTTGAAAAATAACAACCTCAGGTTCGTTGGTCAAGCGCTTAAGAC 2000
ATTGTTTTCAAAATTTTTCATCAACTGTTTCAAACTTTTGTGACATATTATCTCTCAACTTTTATTGTTGAGTCCAGCAACCACTTCCCCAATTCTG

449 T K V . 452

2001 ACCGCTTTTCACGGCGGTAAACGGGTTGAAATCCCGTACCGACTATGCTATGTTGGGTTGGAACACTTGATGAAAACTTTA 2044
TCCGGAAGTCCCGCACTTGTGCGCAAGCTTAAGGATGCTGATACCATACAACGCCAACCTTGTGAACACTCTTTTGAAT

51489

(SEQ ID NO:11) : AACAGCTGCTGTATTTATCTTACCAAAATCCCTCAAATTAGCTAGTAGCATAGCCTTTTGTACTGGCTAAAAACAGGCTATTCCAATTTCG
 (SEQ ID NO:12) TCTCGAGGAAGAATAAATAGAAATCTTTAAAGGCATTAATCGATCATCTATCGACAACATGACCGATTTTGTCCGATAAAGTTTAAGTC

(SEQ ID NO:10)

[illegible]

M H K I L L I E D D O V I R C 15

201 CAGATTGGGAAAAATGCTCTCTCAATGGCGATTNNAAGTGGTCTCTGTACAGACCTTTATGGAGTTTTCAGTCTATTGTTCAGTGGGAACCTCATCTGG 300
CTCTAACCTCTTACGAGACATTAAGCTTAAGTCTCAGCAGGACCATCTCTCTGAAATAGCTTCAAACTCAGATAAACAAGTCAGCTTGGAGTAGACC

16 0 : C K M L S E W G F X V V L V E D F M E V L S L F V O S E P H L V 49

301 TCTTCATGGATATTGGTTTGGCTTGTGTTAATGGTTATCACTGGTGTGAGGAATCCGCAAGATTTCGAAGTACCTATCATGTTTCTTTCTTCGAGACA 400
AGGATTAATTATAGCAAACTGGCAAGAAATTACCAATAGTCAGCAGAGTCTTTAGCGTTCTAAGGTTTCATGGATAGTACAAAGAAGAGAGCTCTC

50 L M D I G L P L F N G Y H W C Q E I R K I S X V P I M F L S S R D 82

431 CCAGGCTATGGATATTGTCATGCGCAATCAATATGGGGGCGGATGACTTGTGACCAAGCCTTTTGACAGCAGGTTCTTTAGCTAAGGTTTCAGGCTTC 508
GGTGGCAATAGCTATAACAGTACGGCTAGTTAATACCTGGCTACTGAACACTGCTTGGGAAAAGTGGTGGTCAAGAAAATCGATTCCAAAGTCCCGAAC

8) O A M D I V M A : N M C A D D F V T X P F D Q O V L L A X V Q G L 115

501 TTGGCTCTTCTTATGAGTTTGGCGGTGATGAGAGTTTCTGGAATATGCTCGTGTATCTTCAATACCAAATCCATGGATTACATTATCAAGGCAAG 600
ACCGGACGACGATCTGCAAGCGGCTGCTGCAAGCACTTATCTACCGAGCAATACGAGTTATGGTTTAGGTACTTAAATGTAATAGTTCGGTTC

116 L R R S Y E F C R D E S L L E Y A G V I L N T X S M D L H Y Q G O V 145

601 TCTTGAATTTGACCAAGATGAATTCAGATTTACGGCTGTTATTGAGCATGCAGGCAACATCTGTAGCACGTGACGACCTGATGCGGGAACTTTGCA
AGAACTTAACTCTCTTACTTAAGCTGTAATAATGCGCAATAAATCTGACCTCGGTTGAGCATCTGTGCACTGCTGGACTACGCCCTTGAACCTT

150 L N L T K N E F C : L R V L F E H A G W I V A R D D L M R E L W N 18

[illegible]

18) S D F F I D D N T L S V N V A R L R K K L E E O G L V G F I E T R 21

80: AAAGGAATACGGTACGGATTGAAGCATGCTGATTGGAACAAATTTTCTAGCCTATCTGCCCTCCCGTAAGTGGTCTTTTATCTATCTGCTTTCTTTC
TTTCTTATGCTTATGCTTAATCTGTAAGCACTAACTTTTGTAAAAACATCGGATAGACGCGAGGGCATCAGGCAGAAAAATAGATAGACCAAAAGAAAC 90

216 K C I G Y G L K H A . 22

901 GCATTCTGCTTACTCTTTCAGTCTTTATTTGCCAGTCTAGGAATTTACTTCTCTACTTTTCTTCTGTGTGTGCTTTGTAAACCATCTATTTTCA
CGTAAGACAGCAATGACAACTTAAAAATAAAGCTTAGATCTTAAATGCAAGGACATGAAAAAGAGAACACAGCAAGAAACATTTGTTAGATAAAAAAGT

gcp1493

(SEQ ID NO:14) : TAAAGACACTGGAAACCAACACCTTCCGCACTTTAGGTAAAGAAAGCTGGTATGGCAACCTTTGTGATTGACTTTTCAAAGGAACCTTAGCAACGGTG 100
(SEQ ID NO:15) : ATTTCTGTGACCTTGGTGGTGTGGAAAGCGTAAATCCATTCTTTCCACCATACCGTTGGAAACACTAACTGAAAGATTTCCTTGGCATCGTTGGCGAC
(SEQ ID NO:13) : K D T G C T T H T F R I L G K K A G N A T F V I D F F K C T L A T L 33

101 CTTCCGATTATTTTCATCTACAAAGCGTTTCTCTCTCATCTTGGACTTTGGCTGTATCGGCCATACCTTCCCTATCTTTGCAGGATTTAAAGGTG 200
GAAGGCTAATAAAAAGTAGATGTTCCGCAAGAGGAGAGTAGAAACCTGAAACCGACATAGCCGCTATGGAAGGGATAGAAAAGTCTCTAAATTTCCAC
34 L P I I F H L Q G V S P L I F G L L A V I G H T F P I F A G F K G G 67

201 GTAAGGCTGTCCCAACCACTGCTGGAGTGATTTCGGATTTCGGCTATCTTCTGTCTCTACCTTGGGATTATCTTCTTTGGACTCTCATATCTTGGCAG 300
CATTCGGACAGCGTTGGTCACGACCTCACTAAAAGCCTAAACCGCGATAGAAAGACAGAGATGGAAACCTAATAGAAAGAACTCAGAGATATAGAACCTTC
68 K A V A T S A G V : F G P A P I F C L Y L A I I F F G L S Y L G S 100

301 TATGATTTCACTGTCTAGTGTCAAGCATCGATCGCGGCTGTTA 344
ATACTAAAGTCACAGATCACAGTGTCTAGCTAGCGCGACAAAT
101 M I S L S S V T A S I A A V 114

gsp1507

(SEQ ID NO:17) : CTAAAGGTAAATCGAATGAAAAGTATAAAATTAATGCTCTATCTTACATGGCAATTCCTGCTTGAATATTATTTCCCATCTAACTGGAACTATG 100
(SEQ ID NO:18) GATTTCATTAACTTACTTTTCATATTTAAATTTACGAGATAGAAATGTAACCTTTAAGCAGACAACTTATAATAAAAAGCGTAGGATTGACCTTGGATAC
(SEQ ID NO:16) : M K S I K L M A L S Y N G I R V L N I I F P I L T G T Y V 29

101 TCGCGCTGTCTTGGACCGAACTGACTATGGTACTTCAACTCACTGCACTATTTGTCATTTTCTTGGCTTTGCAACTTATGCTGTCTATAACTA 200
AGCGCGCACAGAACTGCTTCACTGTATACCAATGAAGTTCAGTCACTGTGATAAAAACAGTAAAAAGAAACGGGAAACGTTGAATACCAAGATATTGAT
30 A R V L D R T D Y G Y F M S V D T I L S F F L P F A T Y G V Y H Y 62

201 CGGTTTAAGGGCTATCAGTAATGTCAAGGATAACAAAAAGATCTTAACAGAACCTTTCTAGTCTTTTATTGTCGATCGCTTGTACGATTTTGACC 300
GCCAAATTCGGATAGTCATTACAGTCTCTATTGTTTTCTAGAAATGTCTTGGAAAAGATCAGAAAAAATAAACACGTAGCGGAACATGCTAAAACTGG
63 G L R A I S N V R D N K K D L N R T Y S S L F Y L C I A C T I L T 95

301 ACTGCTGTCTATATCCTAGCCTATCTCTCTCTCTTTACTGATAATCCAATCCTCAAAAAGGTCTACCTTGTATGGGGATTCAACTCATTGCCCCAGATT 400
TGACCAAGATATAGGATCGGATAGGAGAGAGAAATGACTATTAGGTTAGCAGTTTTCCAGATGGAAACAATACCCCTAAGTTGAGTAACGGGTCTAAA
96 T A V Y I L A Y P L F F T D N P I V K R V Y L V M C I O L I A O I F 129

401 TTTCAATCGAATGGGTCAATGAAGCTCTGGAAAATTACAGTTCTCTCTTTACAAAACCTCC 460
AAAGTTAGCTTACCCAGTTACTTCGAGACCTTTTAAATGTCAAAGAGAAAATGTTTTGAGC
130 S I E W V N E A L E N Y S F S F T K L 148

9ep151:

SEQ ID NO: 20) : CCGCCATTACCGTGAATGATTCACGTATGTAATGATTTTATCGACAACTGAGAGCAGGACGAGAAATGATGTTTGTGACGAGTTCCTATACA 100
CGACCGTAATGGCACTACCTAAAGTGCATACCTACTAAAAATACCTGTTCCAGCTCTCGTCTGCTCTTACATACAAAAACACTGCTCAAGCATATGT

SEQ ID NO: 21) 101 GCGAGTAGGCAATGCAGATTCAAAAAAGTTTAAAGGGCAGTCTCCCTATGGCAAGCTGTATCTAGTGGCAACGCGGATTGGCAATCTAGATGATGACT 200
CCCTCATCGGTACGTCTAAGTTTCAAAAATTTCCCGCTCAGAGGGATACCGTTCCGACATAGATCAACGTTCCCGCTAACCGTTAGATCTACTATACTGA

SEQ ID NO: 19) 1 M O I O K S F X G O S P Y G K L Y L V A T P I G M L D D M Y 30

201 TTTCTGCTATCCAGACCTTGAAGAAGTGGACTGGATTGCTGCTGAGGATACCGCAATACAGGGCTTTTCTCAAGCATTTTGACATTTCCACCAAGC 300
AAAGCAGGATAGGTCTGGAACTTTCTCAGCTGACCTAACGACGACTCTATGCGGGTTATGTTCCGAAAAACGAGTTCTGTAATAAGTGTAAAGGTGGTTCC

31 F R A I O T L K E V D W I A A E D T R N T G L L L K H F D I S T K Q 64

301 AGATCAGTTTTCATGAGCACAATGCAAGGAAAAAATTCCTGATTGATTGGTTCTTGAAGCAGGGCAAGTATTGCTCAGGTCTCTGATGCGCGTTT 400
TCTAGTCAAAATGCTCTGTTACGTTTCTTTTAAAGGACTAACTAACCAAGAACTTTCTGTCGCTTTCATAACGAGTCCAGAGCTACGGGCAAA

65 I S F H E H N A R E K : P D L I G F L K A G O S I A Q V S D A G L 97

401 GCGTAGCATTTGAGACCTGCTCATGATTTTAAAGGAGCTATTGAGGAGAAATTCAGTTGTGACTGTTCCAGGTACCTCTGAGCAATTTCTGCC 500
CGGATCGTAAAGTCTGGGACGAGTACTAAATCAATTCGTGCACTACTCTTTTAAAGTCAACACTGACAGGTCTGAGGAGCTGCTTAAAGACGG

98 P S I S D P G H D L V K A A I E E E I A V V T V P C T S A G I S A 130

501 TGCATTGCCAGTGGTTAGCGCCACAGCCATATCTTTACGGTTTTTACCGAGAAATCAGGTCAACAGAGCAATTTTGGCTCAAAAAAGATT 600
AACTAACGGTCACCAAAATCGCGGTGTCGGTCTATAGAAAAATGCCAAAAATGGCTCTTTAGTCCAGTTGTCTTCTGTTAAAAAACGAGATTTTCTAA

131 L I A S G L A P C P H I F Y G F L P R K S G Q Q K Q F F G S K E D Y 164

601 ATCCTGAACACAGATTATGAATCACTCATCTGTAGGAGACACGTTGGAAAAATGTTAGAACTCTACGGTGACCGCTCGGTTGTTTGTGTCAG 700
TAGGACTTTGTGCTAAAAAATACTTAGTGGAGTAGCACATCTCTGTGCAACCTTTTATACAACTCTCAGATGCCACTGGGAGCCAAACAAACCAATC

165 P E T Q I F Y E S P H R V A D T L E N M L E V Y G D R S V V L V R 197

701 GSAATGACCAAAATCTATGAAGAATACCAAGAGGTACAAATTTCTGAATGCTGGAAAGCATCTCTGAAACGCTCTCAAGGGTGAATGCTCTGATT 800
CCTTAAGTGGTTTATGATACTCTTATGCTTTCTCATGTTAAAGACTTAACGACCTTTCTGAGAGACTTTGAGAGAGTTCCCACTTACAGAAAGACTAA

198 E L T K I Y E E Y O R C T I S E L L E S I S E T S L K G E C L L I 230

801 GTTGAAGGTGCCAGCAAGCTGTGGAGGAAAGCATGAGGAAGCTGTTCTTAGAAATCCAAGCCGATCCAGCAAGGCATGAAGAAAAATCAAGCTA 900
CAACTTCCAGGTGCTTTCCACACTCTTTCTCTACTCTCTGAAACAGAACTTTAGGTTCCGGCATAGGTGTTCCGTACTCTTTTATGTTGAT

231 V E G A S K G V E E K D E E D L F L E I O A R I O Q G M K K N O A I 264

901 TTAAGGAAATAGCTAAGATTACCAGTGAATAAGAGTCACTCTACGCTGCCCTACCAGACTCGGAAGAAAAACAATAAGGGAGACAGGATGTAATAA 1000
AATTCCTTTATCGATTCTAAATGGTCAGCTTATCTCAGTTGAGATCGGACGGATGGTGCTGACCTTCTTTTGTATTTCCTCTGCTCAATTAT

265 K E I A K I Y O W N K S O L Y A A Y H D M E E K O * 290

9ep1518

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(SEQ ID NO: 23) 1 ATGGCTTGCTTAAAAAAGGTGGCAATGCTCTTAAAGTGCAAGTTATTGCGCTGTAGCATATAAATCTATTCTACATATTTTAAAGCTTCTACGAG 100
(SEQ ID NO: 24) TACCGAACCAATTTTCCACCCTTACGAGAAATTCAGTTCAATAACGGGACATCGTATATTAGATAAAGGATGTATAAAAAATTCGAAGATGCTC

101 TTAATTGAAAGCTTTAGCTTGTGTTATATAGATTTATGCGATAAAAAATATGAAAAATCTCTCAGGATTTGGAGTGACGTTAAAGCAATTCATACC 200
AATTAACTTTGCAATTCGAACACCATATTATCTAAATACCTATTTTATACTTTTATAGAGAGTCTTAAACCTTCACTGCAATTTGCTTTAACTATCG

(SEQ ID NO: 22) 1 M D K K Y E X I S Q D L G V T L K Q I D T 21

201 GTTCTAAGTTTGACAGCTGAAGGGGGGACTATTCCCTTTATCGCGCTTATCGCAAGGACATGACTGGTAGTCTGGATGAGGTGGCGATTAAAGCTATTA 300
CAAGATTCAAAGTGTGACTTCCCGCTGATAAGGGAATAGCGCGCAATAGCGTTCTGTACTGACCATCAGACCTACTCCACCGCTAATTCGGATAAT

22 V L S L T A E C A T I P F I A R Y R K D M T G S L D E V A I K A I I 55

301 TTGATTTGGATAAAAGCTGACAAATCTCAATGACCGTAAGGAAGCTGTCTTAGCTAAGATTCAAGAACAGGTAAAGTTGACCAAGGAAATGGAAGAAC 400
AACTAACTCTTTTACAGCTGTTTAGAGTTACTGGCATTCCTTCGACAGAAATCGATTCTAAGTCTTGTTCCTTCACTGCTGCTTAACTCTTCTG

56 D L D K S L T N L W D R K E A V L A K I Q E O G K L T K E L E E A 88

401 TATCTTAGTTGCGGAAAAATTAGCAGACGTTGAAGAACTCTATCTCTTATAAGGAAAAAGCGTCTACCAAGGCAACCTTGGCGGTGAAGCTGAGCTC 500
ATAGAAATCAACCGCTTTTAACTCTCTGCAACTCTTTCAGATAGAAAGAAATATCTCTTTCGCACTGATGTTCCGTTGTAAGCGGCACTTCGACCTGAG

89 : L V A E K L A D V E E L Y L P Y K E K R R T K A T I A R E A G L 121

501 TTTCTCTTCTCTGCTGTTGATTTTGCAGAAATATAGTTGACTTAGAGAAAGAGCTGAAAAATTCCTCTGTGAAGGATTTGGACTGGCAAGGAGGCTTGA 600
AAGGAGAAACGAGCAAACTAAAGCTCTTATATCAACTGAATCTCTTCTCTGAGCTTTTCAAGCAGACACTTCTTAAACGCTGAGCGCTTCTCTGGAAT

122 F P L A R L I L Q N : V D L E K E A E K F V C E G F A T G K E A L T 155

601 CCGGTGCACTTGATATTGCTGCAAGCTTATCGGAAGATGTGACCTTGCCTTCTATGACTTATCAGGAAGTGCTGAGCACTCTAACTCACTTCTCA 700
GGCAGCTCAACTATAAACCAAGCTTCGAATAGCTTCTACACTGGAACGCAAGATACTGAATAGCTCTTCAAGCTCTGTGAGATTGAGTGAAAGCT

156 G A V D I L V E A L S E D V T L R S M T Y Q E V L R H S K L T S Q 188

701 AGCCAGGATGAAAGCTTGTATGAAAGCAGGTTTTTTCAGATTTATTATGATTTTTCAGAGACAGTTGGAAGTATGCAAGGCTATCGTACCTTGGCTCTC 800
TCGGTCTCTACTTTTCAGAACTACTTTTCGTCGAAAGGTCTAAATAAATACTAAAAAGTCTCTGCAACCTTGATACGTTCCGATAGCATGGAAACGAGAG

189 A K D E S L D E K Q V F O : Y Y D F S E T V G T H Q G Y R T L A L 221

801 AATCGTGGGAGAACTTGGTGTCTGAAGATCGGTTTTGAACATGCGACGACCGTATTCTTGCCTTCTTGTCTACTGTTTCAAGGTGAAAAATGCTT 900
TTAGCACCTCTTTTGAACCAAGAACTTCTAGCCAAACTTGTACGCTGCTTGGCATAAAGACGGAAGAAACGATGAGCAAGTTCCACTTTTACGAA

222 N R G E K L G V L K : G F E H A T D R I L A F F A T R F K V K N A Y 255

901 ATATTGATGAAGTTGTTTCAGCAATCGTTAAGAAAAAGGTCTTGCCTGCTATTGAGCGTCTATTTCGGACAGAAATTAAGTGAAGAAAGCTGAAGAGGAGC 1000
TATAACTACTTCAACCAAGTCTTAGGCAATCTTTTTCCAGAACGAGGATACTCGCAGCATAAGCCTGTCTTAATTGACTTTTCAAGCTTCTCCCTCG

256 : D E V V O O S V K K K V L P A I E R R I R T E L T E K A E G A 288

1001 TATCCAACCTTTTCTGACAACTGCGCAATCTCTCTGCTGCTCTCACTGAAAGGGGCGCTGCTTCTGGAATTTGACCCAGCCTTCTGTACAGGTGCG 1100
ATAGGTTGAAAAAGACTTTAGACGCTTAGAGGAGAACCAAGAGGTGACTTTCCCGCGCACCAAGAACCTAACTGGGTGGAAGACATGTCACCG

289 : Q L F S D N L R N L L L V A P L K G R V V L G F D P A F R T G A 321

1101 AAGTTAGCTGTCTGGATGCAACAGGAAAAATGCTGACAACTCAGGTTATTTATCTGTTAAACAGCATCAGCTCTGTAATCGAAGAAAGCCAAAGAAAG 1200
TTCAATCGACAGCACTAGCTTGTCTTTTACGACTGTTGAGTCCAATAAATAGGACAAATTTGGTGTAGTGTGAGCAGTTAGCTTCTTGGTTCTTTC

322 K L A V V D A T G K M L T T Q V : Y P V K P A S A R Q I E E A K K D 355

1201 ATTTAGCAGATTAAATGCTCAATCGGTGTAGAGATTATTGCAATGGAATGGAACGGCCAGTCTGAAAGTGAAGCTTTTGTAGCGGAAGTTCTGAA 1300
TAAATCTCTAAATTAACAGTTATGCCACATCTCTAATAACGGTAAGCTTACCTTGGCGGTGAGCACTTCACTTGAAGAAACATCGCTTCAAGACTT

356 L A D L I G Q Y G V E I I A I G N G T A S R E S E A F V A E V L K 388

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1301 AGATTTCCCTCAAGTCAGCTATGTTATCTTAAATCAAAAGTGGTCTTTCTTATTTCTGCCAGCCAACTTGCTGTCAGGAGTTCCAGACTTGACCTT 1400
TCTAAAGCGACTTCAGTCCATACAAATAGCAATTACTTTCCACGAGGAGACAGATAAGACGGTCTGTTGAAAGGAGCTCTTCAAAGGTCTCAACTGGCA 1400

1389 D F P E V S Y V I V M E S G A S V Y S A S E L A R O E F P D L T V 421

1401 GAAAAAGCTCTGCCATTCTATGCGCCCTCTCTTGCAGATCTCTTCCGGAATTGGTCAAAATCGATCTTAAGTCAATTTGGTGTGGTCAATACCAAC 1500
CTTTTCCGAGACGGTAAAGATAGCGGGCAGCAACCTTCTAGCAGAACGGCTTAACCACTTTAGCTAAGATTCACTTAACCAAGCCAGTTATGGTTC 1500

1422 E K R S A I S I A R R L O D P L A E L V K I D P K S I G V G Q Y Q H 455

1501 ACCATGTCAGTCAGAACAACTATCTGAGACTCTGCACTTTCTTGTGGATACAGTCTTAACCAAGTTGGTCAATGTCATACAGCTAGCCGCTCTT 1600
TCTACAGTCACTCTTCTTATAGACTCTCAGACCTGAAACACAGCTATGTCACCAATTTGGTCAACCAAGTTACAGTTATGTCGATCGGTGGAGA 1600

1456 D V S Q K K L S E S L D F V V D T V V M Q V G V N V M T A S P A L 488

1601 TCTTTCAAGCTAGTGGACTCAACAACTATCTCTGAAATATTTGTCAAAATACCGGAGCAAGCAAGCAAAATCACTTCAAGCCGCTCAATCAAGAAA 1700
AGAAAGTGTCCATCCACTGAGTGTCTTATAGAGACTTTTATACAGTTTATGGGCTCTTCTTCTTTTATGTAAGTGGCGGCTTATGTTCTT 1700

1489 L S H V A G L N K T I S E N I V K Y R E E E G K I T S R A Q I K K 521

1701 GTTCTCTGCTGGGAGCCAAAGGCTTTGAGCAGGCTGCTGCTTTCTTCTATCTCTGAAAGTAGCAATATCTTGATAATACAGGAGTTCAAGCAGAG 1799
CAAGGAGCAGACCTCTGGTTCCGCAAACTCTGCGACGACCAAGGAAGCATAGGCACTTTCATGTTATAGGAAGTATTATGTCCTCAAGTGGGCTCT 1799

1522 V P R L C A R A F E Q A A G F L R I P E S S N I L D N T G V H P E 554

gcp1846

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(SEQ ID NO: 26) 1 TACTGGGCGAAGCGTTTCTTACCTGTTCTGAATGTGAAGTCTTCTGAAAATGCTGAAGTTAAGATTTTCAGAGCACTCAACGCAAGCCAGNATCTCT 100
(SEQ ID NO: 27) 1 ATGACCCCGTTCCCAAGAAATGGGACAGACTTACACTTCGAGAAAGAACTTTACCACTTCAATTCTAAAAGTTCTGAGTTGCTTCGCTGTATAGCGC 100
(SEQ ID NO: 25) 1 T G A R V S Y P V L E V R V F L E N G E V K I P R A L N E A X I R 133

101 AGGCTCTGATCGAACCATGCTGGCAGATATTGTAAATAATGCTTCTCTTCAACGTTTCTGTCAGAGCGGCTAAGAGTTTCSACACCGACTGCTAGTA 200
TCCAGACTAGCTTGGTACCACCGTCTATAACATTATTACGCAAGGGAACCTTCGAAAGCACTTCTGCCGATTTGCAAGCTGTGCTGACCATCAT 200
34 R S D R T M V A D I V I N G V P F E R F R C D G L T V S T P T G S T 67

201 CTGCTATAACAAGTCTCTTGGCGTCTGTTTACACCTTACCATTAAGCTTTGCAATTAACGGAGATTGCCAGCTTAATAATGCTGTCTATGAAAC 300
GACGATATTGTTTCAAGAACCGCCACGACAAATGTGGATGCTAACTTCGAAAGCTTAATTGCTCTAACCGTGGGAATTATTAGCACAGATAGCTTC 300
68 A Y N K S L G C A V L N P T I E A L O L T E I A S L N H R V Y R ? 100

301 ATTGGGCTCTTCCATTATTGTCCTAAGAAGGATAAGATTGAACCTTATCCACAAGAAAGGATTAATCACTATTTCGCTTCACAAATAGCTTTATTCT 400
TAACCCGAGAAAGTAATAACACCGATTCTCTTATTCTAACTTGAATAAGGTTGTTCTTCTGTAATAGTATGATAAGCCAACTGTTATCGCAATAAGA 400
101 L G S S I I V P K K D K I E L I P T R N D Y N T I S V D N S V Y S 133

401 TTCCTAATATTGAGCGTATTGAGTATCAATCCACCATCATAAGATTCACTTTCTGCGACTCTAGCCATACCACTTTCTGCAACCGCTTTAAGGATG 500
AAGGCATTATAACTCGCATACTCATAGTTAGCTGGTAGTATCTAAGTGAAAGAGCGCTGAGGATCGGTATGGTCAAGAGCTTTGGCACAATTCCTAC 500
134 F R N I E R I E Y Q : D N K I N P V A T P S R T S P M N R V K D A 167

501 CCTTTATCGGTGAAGTGCATGAATGAGCTTTGAATTTATCGGAGATGAACATGTCAAGCTTAAGACCTTTTAAAAAA 578
GGAATAGCCACTCCACTACTTACTCCAACTTAAATAGCTCTACTTGTACAGTTCCAAATTCGGAAAAATTTTTT 578
168 F I G E V D E . 175

gcp1331

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(SEQ ID NO: 29): CCCTCTAAAGAAACCTACTGACAGTGCATAGATGGGAAGTACTATTAATTCCTTTATCCGAGAGATGCTTGTCCGCTGGCAATATATACCTGCT 100
(SEQ ID NO: 30) CCGAGATTTTCTTGGATGACCTCTCACTATTATCTCTTCATGATAATAAACTAGGAAATAGGCTCTCTTACCAACAGCGACCTGTATATATGGACCA 100
(SEQ ID NO: 28): M V V G H O Y I P A 10

201 CCACACAGGGGTTACGATTCCTCTCTCCAGCAATAGAGATTCTCTTAGACCAGATTGCTTTATTTTGGTCAAGATGCTCTTACAAGAATTTC 200
GGTGTGTTCCCTCCAAATGCTAACCCAGGAAGAGCTTTATTTCTTAACGAGAAATCTGGTCTAACCCAAATATAAACCACTCTACCAAGATTTCTTAAAC 200

11 P H K G V T : G P S P R I E I A L E P D W F Y F G O D G V L O E P V 44

201 TTGCCAAGCAAGTTTATGAAGCAAAACCTCTACCAATACCAACAAATCATGCGGAAGATATCATAGCCAGGAGAGAACCACTCTATTATTTTGA 300
AACGTTGCTTCAAAATCTTCTTTTGCAGATGCTTATGCTTCTTGTAGTAGCCCTCTTATATCTATCTGCTTCTCTTCTCTCAGATATAAAACT 300

45 G K Q V L E A K T A T N T W E H N G E E Y D S Q A E R V Y Y F E 77

301 ACATCAGCGTAGTTATCATACTTTAAAACTCTTGGATTATTAAGAGCGTTATTCCTATTATTTACAGAAAGATGCTGCTTTGATTCCTCAATCAAC 400
TCTAGTCCCATCAATAGTATGAAATTTTCAACCACTAAATATCTCTCCAAATAACCAATAATAATGCTTCTTACCAACCACTAAGAGCGTAGTTG 400

78 D O R S Y N T L K T G M I Y E E G Y W Y Y L O K D G O F D S R I N 110

401 ACATTGACGCTTGGAGAGCTAGCACCTGCTGGTTAAGGATTACCTCTTACGTATGATGAAGAGAAGCTAAAGCAGCTCCATGCTACTATCTAGATC 500
TCTAATGCCAAGCTTCTGATCTGTCACCAACCAATCTCTAATGGGAGATGCTACTACTCTCTTCTGATTTTCTGAGGTACCATGATAGATCTAG 500

211 R L T V G E L A R G W V K D Y P L T Y D E E K L K A A P W Y Y L D P 144

501 CAGCAACTGGCTGCCAAACCTTCCGCAACAAATGCTACTACTCTCTCTCATCAGGAGCTATGCTAAGTGGCTGCTATCAGATGCTTTAAGTGGTACTA 600
GTGCTTGACCGACCTTTTGAACCTTCTTACCAATGATGGAGGCAAGTAGTCTCTGATACCAATGACCGACCATAGTTCTACCAATGAAACCATGAT 600

145 A T C W O N L G N K W Y Y L R S S G A M V T G W Y O D G L T W Y Y 177

601 CCTAAATGCCAGTAAATGGAGACATGAAGACAGCTTGGTTCCAAGTCAATGGTAACTGGTACTATGCTATGATTGAGGTGCTTACCTGTTAATACCA 700
GGATTACGTTCAATTACCTGTGTACTTCTGTCGAACCAAGTTCACTTACCAATGACCATGATACCGATACTAAGTCCAGCAATCGACAAATTATGGTGT 700

178 L M A G N G D M K T C M F O V N G W Y Y A Y D S G A L A V N T T 210

721 GTAGGTGGTACTACTTAACTATAATGGTGAATGGTTAAGTAATGAAGGCTAAATGTAATCTGTGATGGATACTTAACTTTGTATAATAGGTGATAA 800
CATCCACCAATGATGAATTTGATATTACCACTTACCAATTCATTACTTCCGATTACATTGACACTACCTATGAATTGAACATATTATCCACTATT 800

211 V G G Y Y L N Y N G E W V R 225

99p1861

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(SEQ ID NO: 32) : TTTATGGATATTTATTTAAGAAAAGCCATTTACCAGTTCAGTCCGGATGATACCGAGCTTTCTTAGCAGATAAGTTTCTCAATATTACTCCAAA 100
(SEQ ID NO: 33) : AAAATACCTATAAATATAATTTCTTTGGTAATAAGTGGTCAAGTCAGGCTTACTATGGCTCGACAGAAGATCGTCTATTCAAGAGTTATAATGAAGTTT 100
(SEQ ID NO: 31) : N D I Y I K K A I I N O F S P D D T E L F L A D E F L N I T P R 11

101 ATCGAAGATACCTACGTAAGAAAATTCACATCTGTATTACATGAAGCCAGACTGGGATTTCGAGAGAAGAAAATCCCTCTCAATCATATTACAG 200
TAGCTTTCTATGGATGCAATTTTAACTTGTACACATAAGCTTACTTGGTTCGACCTAAGAGCTTCTCTTTTAGGAGAAGATTTAGTATAATGTC
33 I E E Y L R K K I E N V Y S D E A K T G I F E E E N P F F N N I T D 66

201 ACCGATTTGTGGAGCATCAGTAACGCTGGCTAATCTCTGAAAGAGGAGTTTACGATTTCTCAAAATCTCAAGACCAATGACTTGTATTTGTTCATT 300
TGTAAAGCAACCTCTGTAGTCATTGCGACCGATTAGAGACCTTCTCTCAAAATCGTAAGAGCTTTTAGAGTTCTGGTTACTGCACTAAAAGCAAGTTAA
67 D L L E T S V T L A N L W R E E F S I S E N L K T N D L I F V Q F 99

301 TTTAAGAGAAGGTGACAACTTTCCCTTTCTGCAATTTGCTCTCGGAGACCTTCAACCACTCGGAGGAGAAGTTGATATATCAATCAAGCTGACT 400
AAGATTTCTTCACATCTTGTAAAGCGAAGAGAGCTTAAAGGAGAGCTTCTGCAACTGGGTGGAGCTTCTCTTCAACTATTAGCTTACTTCGAGTGA
100 S K E O V E N F A F L R I A L R E T L T N L G G E V D N P I K L T 132

401 CAGAATAACCTGCTGGATTTGGAACGGGTGCTGACGAGGCTTGGTGTCAATCTTCAGAGTCGCAAGTATCACCTGATCAAAAACCAATCAAGTACA 500
GTTTATTTGAGCGGACCTAAACCTTGGCCAGCTGCTCGGAGACCAAGTATAGAGTCTCAGCTTATATGAGTAACTTTTGTCTAGTTGATGT
133 Q N N L P C F G T C A D E A L V V N L O S R K Y N L I E R R I K Y N 166

501 ACCGGACTTTTGACTATTTTCAGATAATCTTCTGCTGCTGCTCTCAAGATTTCTCTAAAAATCTATCAAGGAATCGAAAAACAGCCGAGAG 600
TGGCTTCAAAAACTTGATAAAAAAGTCTATTAGAGAGACGAGAGAGGATTTCTAAGAGGAATTTTAGATAGTCTCTTGAAGCTTTTGTGCGGTCTC
167 C T F L N Y F S D N L L A V A P K I S P K K S I K E L E K T A Q R 199

601 AATGCTGAATCTTTTACACAGATGATTTTCAATTTCAATCCAGGTCAATCAGCTATTTTCAACCACTAGAGAAGCAATGAATGTCACTGAG 700
TTAAGCACTTAGAAAAATGTTCTTACTAAAAAGTTAAAGTTAGGTTCCAGTTAGTCTGATAAAGTGTGAGATCTCTTCTGTTACTTAACAGTGGACTC
200 I A E S F N T D D F O F O S K V K S A I Y N N L E E S N E L S P E 232

701 AAATGGCTAATGACTTTTGAACAATCTCAGGCTGCTTTGAGCTTTATTGACCAACTCAGAGAAGCGGTACCAGAACCTGTTCAATTCATGAAA 800
TTTAAAGGATTAAGAAAACTGTTTGTAGACTCGGAGCAAACTGAAAATACTGTTCACTGTTTGGCATGCTTCTTCTGTTACTTAACAGTGGACTC
233 K L A N D L F D N N L T A R L S F I D O V R E A V P E P V O F D E I 266

801 TTTATGCGAGTCGCAATTAAGAAAATTTGAAAACCAAAAATCTCTTTATCAATGCAATTCAGCTCATCTTCCCAATAAGCTCTATCAAGACCGGA 900
AAGTACGCTCAGCGGTAATTTCTTAACTTTGCTTTTGAAGCAATAGTTTAACTTAACTCGAGTAGCAAGGTTATTGAGATAGTTCTGCGGCT
267 C A S P O L K K F E N O K L S L S N G I E L I V P N N V Y O D A E 299

901 GTTCTTTGAGTTTATCCAAAACGAAAATGGAACCTACTCTATTTAATCAAAAATATCGAGGATATCAAGTAAATATGTTTAAAGCAATTCGAAGAG 1000
CAGACAACTCAATAGGTTTTGCTTTTACCTTGGATGAGATAGAAATAGTTTATAGCTCTATAGCTTTCATTATTACAAATTTGCTTAAGCTTTCTC
100 S V E F : Q N E N G T Y S I L I K N I E D I O S R - 1125

1001 TGGTTGTAAGTACGAGTCTCTCTTTTGTGCTATAAAGCTTACCGGCTTATCAAGATGTCAAAACAGTCTAGCACTATCAACCCATGGTGGAGAAAAT 1100
ACGAACATGATCTCAGAGAAGAAAACGACCGATATTTGCAATGGCGCAAGTACTTCTACAGTTTGTTCAGTACTGATAGTTGGGTACCAAGCTCTTTTA

gcp1380

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(SEQ ID NO: 35) : AATGTGCTATAATACACAAAAATCTTGTGAGGTTCCATTATGCCAATATTTTCATGATTTTCTGATTGTTGTGTGCTCTATTGTTGATAGTCT 100
(SEQ ID NO: 36) : TTACAGCATATTATGATCTTTTATGAACACTTCCAGGTAATACCTTTATAAAAAGTACTAAAAAGACTTAACAAACACACAGGATTAACCACTATCAG 100
(SEQ ID NO: 34) : H A I F F H I F L I V C V L L L V I V 19

101 ACACCTGAGTACAGTTTATGTGCTTCTTCAGCAGTCCGTGCGCAATTATGAACGCTTTGGGAAATACCAAAAGGTTGCTAATACCGGTTTCATATTGCTT 200
TGTGACTCATGTCAATATACACCAACCACTCTTCAGCCACCGCTAATAACTTCCGAAACCTTTATGCTTTTCCAAAGATTTATGCGCATTAAGTATAAGCGA 200
20 T L S T V Y V V R Q Q S V A I I E R F Q E Y O K V A N S G I H I R L 59

201 TGCCTTTTGGGATTCAGTCTGATTCAGCAGCGGATTCAGTTCGCTTCTTCCAAAGTATATTTGCTGAGACTAAGACCAAGGACAAATGTTCTTCTTAT 300
ACGGAAGACCTTAAGTCACTAAGCTGCTCAAGTCAAGCGCAACAGCTTTCAGTATAACACCAACTCTGATTTCTGTTCTCTGTTACACAGCAATA 300
54 P F C I D S I A A R I Q L R L L O S D I V V E T E T E D N V F V H 86

301 CATGATGTAGCCACTCAGTACCTGTCTAACGAGCAGAGCGTGACAGATGCTTACTATAAACTCATACCTCCAGAACTCAGATTAATCTTATATGCA 400
CTACTTACATCGCTCAGTCTAGCAGCTTGTCTGCTCTGCACTGTCTAGCAATGATATTTCAGTATGCAAGCTTTCAGGTTCTAATTTAGAAATAGCTT 400
67 M N V A T O Y R V N E O S V T D A Y Y K L I R P E S O I X S Y I E 119

401 CATGCTCTTGGCTCTTCTGTTCCAAATTAACCTTGGATGAATTTGTTGCAAAAAGATGAGATTGCTCTTTCAGGTTTCAACCAAGTACGAGAGAAA 500
CTAGGAGAGCGAGAGAGCAAGGTTTAAATGGAACTTCTTAAACAACTCTTCTTACTCTAACCGGAAGTCCAAAGTTGTGCTTCTGCTCTCTTCTT 500
120 D A L R S S V P K L T L D E L F E K K D E I A L E V O N O V A E E H 153

501 TGACCACTTACCGCTACATTATGCTGAAAACTTCAATTACCAAGTCCGACAGATGCAAGTAAAGCAATCTATGAATGAAATCAATGCCGGCGCAAG 600
ACTGCTGAATGCCGATGTAATAGCACTTTGGAATTAATGCTTCCAGCTTGTCTACCTCTTCAATGCTTAGATACTTACTTTAGTTACCGCGCTTCC 600
154 T T Y G Y I I V K T L I T R V E P D A E V K O S H N E I N A A O R 186

601 TAAGCGGCTGCGAGCACAAGAAATGGCGGAGCTGACAAGATTAAATTTGCTGACGCTGAAGCGGAGCAGAAAAAGACCGCTTCTATGTTGTGGGG 700
ATTGCGGCGAGGCTGCTGTTCTTAAACCGCTTCCAGCTGTTCTAATTTTAAACAGTGACCTGCACTTCCGCTTCTGTTTCTTCCGCGCAAGTACCACACCCC 700
187 K R V A A O E L A E A D K I X I V T A A E A E A E K D R L N G V C 219

701 ATTCGCCAACCAAGCTAAGGCGATTGCGATGCAATTGCGCAGCTCTATCACCGAAGTCAAGGAAGCCAAATGTTGCGATGACAGAGAACAAATCATGTTCTA 800
TAAAGGCTTCTGCAATTCGGCTAACCTTACTTAACCGTCTCAGATAGTGGCTTCACTTCTGCTTACAAAGCTACTGTTCTTCTTCTTATGATACAGAT 800
226 I A O C R K A I V D G L A E S I T E L K E A N V G M T E E O I N S I 253

801 TCTCTTACCAACCAAGTATTGGAATACCTTGAATACCTTCTCTTAAAGCAAAATCAAACTATTTTACCAAAATCTCCAAATGTTGCGATGATAT 900
AGGAGAACTGTTGCTGATAAAGCTATGGAAGTATGGAAGCGGAGATTTCTTTAGTTTGGTAGAAAAATGTTTATGAGGTTTACCACACCTACTATA 900
254 L L T N O Y L D T L N T P A S K G N O T I F L P N T P N G V D D I 286

901 CGGTACACAAATCTTCTCAGCCCTTCCGCTGAGAAGAAATATAGACTAATACTCTTCCAAATCTCTTCAAACCTACCTCAGCTGCTCTTCCGCTATA 1000
GGCATGTTTATAGAACAGTCCGGAAGCGGAGCTCTCTTCTTATATCTGATTATGAGAAGCTTTAGAGAAGTTTATGCAAGTCCGAGCAGAACCGCATAT 1000
287 R T O I L S A L R A E K R 300

seq1713

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(SEQ ID NO: 38) : CTTGATATGTTGATATAAATAGGCTTTTATTTCGAAACCTTTTCTGTTTCATTTGCTAAAAAATGCTACAAATAGGAAAGCTTACTATTA 100
(SEQ ID NO: 39) : GGAATATACCACTATTTTATCCCAAAATTAACCTTTTCGAAAGCAACAAAGTTTAACGATTTTTRACCATGTTATTTCTTTCGAAATGATAAT

101 TCTGAATCAGCAGATTTCAGAGAAAGCAATCAATTCATTAATCAATAGGCTTTTTCGAAAGCTGAAAGGCTTCTCTACTAAAGAGCTGATTTTATGCG 200
AGACTTAGTCTGTTAAACCTCTCTTTCTTAAGTAAACTTTAGTTATCCGAAATTAACCTTTTCGACTTCCCAACAGATCAATTTCTGCACTAAAAATAACCC

(SEQ ID NO: 37) : L S I G F I E K L K G L S S K E L I L L C 22

201 AATTATCCTAAGTATCTTTTACCTTTTATCTTTTCTAGTTTACTCTGTTTATATATTATCAGTTTCATTTTACAGCAGACATGAAAGTATTTCTT 300
TTATAGGATTCATAGAAAAATGGGAAATAGAAAAACATCAACATGAGACAAATATATAATAGTCAAACTAAAAATGTTCTCTGACTTTTCATAAGAA

22 I I L S I F L P F Y L F V V V L C L Y I I S L I F T G D N K S I L 35

301 CAGAAATGGGGAGCATCCGATGCTCTTTTCTTAAGTATAGTACTGTTATATCCATTTCCACAAATTCGATGCGCTTTGCGCTTCAGTAG 400
GTTCTTACCCCTCTAGGCTACGACCAAGAAAAAGATCAATATCATGACAAATATAGGTAAGAACGTTTAACTACCCAGAACCCGAAAGTCAATC

36 O R N G E R P M L L L F L S Y S T V I S I L A Q N W M Q L V A S V Q 89

401 GAATGTTCTATTTACTATTTTCTTCTACTATCAGTGGATTTATCCCAATAATCTTTGATTCGATTTTTCAGTTCTGTTCTGTTGCTAGTGTCTT 500
TTATAGGATTAATGATAAAGAAAAAGTCAATAGTCACTAAAAATAGGCTATTTAAGAAAGCTAACATAAACCTGCAAGCAGAACCAATCACAGAA

90 M F L P T I F F L N Y O S I L E N R F P R L I L O F V L P G S V L 122

501 CTCAGCTGCTTTTCCAGTTTAGAACAATTCGAAATTCGAAAGAAATTAACATGCTTTTCTTTCACCAATATCCAGTGTGCGATCAGAACCGGCA 600
CAGTGCAGCAAAACCGTCAATCTTGTAAAGGTTTAACTCTTCTTAAATGATACGAAAGAAAGTGGCTTATAGCTTCACACCGTATGTTTGGCCCT

123 S A A F A S L E N F C I V R K P N Y A F L S P M N Q V M N Q N R A 155

601 GAATGACCTTCTTAACTCAATTAATGGAATTAATGTTTCTGTTATGATGCTTTCTATCTGTTTACAAAGCAAGTTGAATGCTGCA 700
CTTCACTGCAAGAAATAGGATTAATTAATCTTAAATAAACAACAGACATAATAGTAAAGCAAGTACAGAAATGTTGCTGCTCACTTAAGCAACT

156 E V T F P N P N Y Y G I I C C F C I M I A F Y L F T T T K L M W L K 189

701 AAGTATTCGTTGATTCAGGCTTTCTTAACTCTTCTGTTTGAACCTTTACTCAAAATCGAAGTGGCTTTCTGCTATTATCGCTGAGCAATTTCTA 800
TTCAATAGACACTAAGCTCCGAAACAAATAGAGAAACCAACTTCAATAGCTTTAGCTTGACGGAAGGAGCATAATAGGCTCTTAAATAGAT

190 V F C V I A G F V N L F G L N P T O N R T A F P A I I A G A I I Y 222

801 TTTTATTACGACTATTAAGAACTGGAAGCTTTTGGCTTAGTATTGGCTTTTCCGATGCTTTGAGTTTCTCTTCTAGTGAATTTGGAGTTTCA 900
AGAGAAATGCTGATAATTTTGAACCTTCCGAAACCAATCAATACCCGAGAAGCTTAACCAACTCAAGGAGAAAGATCACTAAACCTTCAAGCT

223 L F T T I K N M K A F W L S I C V F A I G L S F L F S S D L G V R 255

901 ATGGTACTTTAGACTCTTCTATGGAAGAACCAATCTATCTGGGATGCTGGGATGGCTTCTTAAAGCAAAATCTTTTGGGTCGAAGGCGCTTGA 1000
TACCAATCAAAATCTGAGAGATACCTTTTCTGTAAGATAGACCTACGACCTACCGGACCAAAATTCCTTTAGGAAAAACCCGACTTCCCGTAAC

256 M G T L D S S N E E R I S I N D A G M A L F K O N P F N G E G F L T 289

1001 CTTATATGCACTCTTATCTCGGATACATGCTCTTATCATCAACATGCCACAGTCTTTATATTGATACGATCTGAGTTACGGAATTTGGGTACCAT 1100
GCATATACGTCAGAAATAGGAGCTATGTACGAGGAATAGTACTTGTACGGGTGTCAGAAATATAACTATGCTAAGACTCAATGCTTAAACCCATGATA

290 Y M N S Y P R I N A P Y N E N A N S L Y I D T I L S Y G I V G T I 322

1101 TTTATTAGTTTCTCTCTGTTGCTCTGTTGCTGTTGATGATGATATGATGAGGAGTCCGGAACCTCCGATTAACGCTTTATCTATCTTTCTCT 1200
AAATATCAAAACAGAGACAAAGGAGCAAGCACTACTACTATATCTAGTCTCTCAGCCCTTTGAGGCTAATAGCCGAAATAGATAGAAAGGAA

323 L L V L S S V A P V R L M M D N S O E S G K R P I I G L Y L S F L 355

1201 ACAGTGGTCTGTCACAGAAATTTCACTTGGCTCTCTCTGATTCAGTCAAGCTTTATTTCTTCTGATGTTATGTCAGTATTCATTTGGCTTTA 1299
TGTCACCAACGACACGCTGCTTAAAACTCAACCGAGAGAGAGCTTAAGTCACTCCGAAATAAAGAACATCAATACAGCTCATAGGTAAACCAAT

356 T V V A V N G I F D L A L F W I O S G F I F L L V M C S I P L A L 388

gsp222

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(SEQ ID NO: 41) : AAGCACTGAACATCTGGCTCGGTACTTCAATTCATGAAAGTATGCGTGAATGAATTCGTGTAACAGTTGTCGCAACGGGTGTTCTGCAAGACCGGTAGA 100
(SEQ ID NO: 42) : TTTCTCACTTTGTAGACCGAGCCATGAAGTTAACTACTTTTCATAGCACTACTTTAAGCACATTGTCACACAGCGTTGCCACAGCACTTCTGGCCCATCT

101 AAGGTGTGCTCCACAGCTAGATCTCTACTAACTACCTGAGACAGTGAACACAGCTCATTACATGGCTTTCATGCTCAATTTGATATGGCAGAA 200
TTTCCAACACCGAGGTGTTCCATCTAGACCATGATTGATGCGCACTGTCTCACTTTGTTGAGTAAAGTGTACCGAACTAGCACTAAAATATACCGTCTT

201 AAGTTGAATTCGCAAAACAAAATCCACTGCTTTGCAACCAACTCAAGCATCTGCTTTGCTGATTGGGATCTTCGCGTGAATGCAATGTTCTGTACAA 300
TGTCAACTTAACGGTTTGTGTTAGGTGCAGCAAACTTGTGCTGAGTCCGTAGAGCAAAACCACTAACTCTAGAAAGCGGCACTTAGCTAACAGCATGTT

301 CAGATTCACTGCTTTCTCACTCGAGCGCTTCAAGCCCAATTTCAAGATCAAGATGAATTCGATACACCTCCATTTTCAAAAATCOTTAAGTAAA 400
ATCTAAGTCAGCAAGAGCTCAGCTCCCAAACTCCGGGTTAAAGTGTCTACTTCTACTTAACCTATGTGGAGGTAAAAGTTTATTAGCAATTCATTT

(SEQ ID NO: 40) : M 1

401 TCAATGTAAAGAAAATACAGAACTGTTTTCGAGAACTGCGAGCGCTACTCTGAGTCTCATCGAGAGAGCTGTTTGGTCTCTGTCATTGCACTTAT 500
ACTTACATTTTCTTATGCTTGAACAAAAGCTCTTCAAGCTCTCGATTCAGACTCAGCAGTACTCTCTCCACCAAGCCAGAGACAGTAACTGCAATA

2 N V K E N T E L V F R E V A E A S L S A H R E S G S V S V I A V I 34

501 CAAGTATGTAGATGTACCGACAGCGGAAGCTTCTTCTGGCTAGGTGTTCAATCATATCGGTGAAAATCGGTAGATAAGTTTTCGCAAAAATATCAAGCT 600
GTTCAATCATCTACATGGCTGTGCTTTCGCAACGAAGCGCATCCCAAGTAGTATAGGCCACTTTAGCACATCTATTCAAGACCTTTTATAGTTCGA

35 K Y V D V P T A E A L L P L C V N H I G E W R V D K F L E X Y E A 67

601 TTAAGATCGAGATGTCACTTGGCATTTGATTCGTACTTTCGCAAGACGTAAGGTGAAGATGTCATTCAATACGTTGATTATTTCCATGCAATGGACT 700
AATTTTCTAGCTGTACACTGAACCGTAACTAACCATGGAAGCTTCTGCAATTCACATTTCTACAGTAAGTTATGCAACTAATAAGGTAGCTAACTGA

68 L K D R D V T W N L : G T L O R R K V K D V I O Y V D Y P H A L D S 101

701 CAGTAAAGCTACCGCGGAAATCAAAAAGAAAGTGACCGAGTCATCAAGTGTTCCTTCAAGTAAATATTTCTAAAGAGAAAGCAAAACCGGTTTTC 800
GTCATTTGATGCTGCTGCTTAAAGTTTTTCTCACTGGCTCAGTAGTTTCAAAAGGAAGTTCAATTTATAAGATTTCTTCTTCTGTTGTCAAAAG

100 V K L A G E : C K R S D R V I K C F L O V N I S K E E S K N G Y S 134

801 GAGAGAGGAAGCTGCTGGAATCTTCCAGAGTTAGCTCAGACTAGATAAGATTGAATATGTTGCTTTAATGACCATGCCACTTTTGAGGCTAGCAGTCAG 900
CTGCTGCTTCAAGCACTTTAGAAAGCTGTCAATCGGTGTGATCTATTCTAATTTATACAAACCAATTAAGTGTACCTGCAAAAGCTCCCATGTCCTCTC

135 P E E L L E : L P E L A R L D K I E Y V G L M T H A P P E A S E E 167

901 CAGTTCAAGAGATTTTCAAGCGGCTCAAGATTTCAAGAGAGAAATTCAGAGAGAAACAAATTCAAATATGCTTTAGAGCACTGCGCGCGCTTAC 999
GTCAACTTTCTTAAAGTTCCGCGGCTTCAAAATGTTCTCTTAAAGTTCTCTTTGTTTAAAGTTTATACGGAATCTGCTGTGACCGCGCGCAATG

160 C L K E : F R A A C D L O R E I O E K Q : P N H P L E N T G G R Y 200

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(SEQ ID NO: 44):	GTACTCCGCTCCACTTTAGCCAGTAAGTTATTATTATTTTAAATGACGCCAACATTTCTCTGCTGCAATTCAGATTTTGTAGTACTTCTGTAATTTCT	100
(SEQ ID NO: 45):	CATGAGGGTCAGGTGAAAAATCGTCACTTCAAAATAATAATGAAAAATAGTCTGGTGTTAAAGAACAGAACTTAGTCTAAAAACCATCCATCAAAACCATTAAGA	
(SEQ ID NO: 43):	T P S P L L A V S L L F T T F N O P O F L V L N Q I L V G S L V I L	33
101	ACTTATTGCATATATAGTGTAAAAATCCCATTTCTTTATACAACTGCTAGCTCTATTTTATTAGTGTTCATGATGAGATGGAAGATGCCAAGAACT	200
34	TCATAACGCTATATATCAACATTTTtagggtaaaagaaTATCTTACCATGCGCATAAAAATAATCACACTACTACTCTACTCTTCTAGCTCTCTCTCA	66
201	ATGGGTGCTTCACCTTTTATACATATGATGAAGGTTATCATTCATTTTACCGGCTCTCTCTCTGTTATTCCTTTAACTTTAACTCTTTATTAA	300
67	TACCCACGGAAGTGGAAAAATATGATACTACTTCCAAATAGTAAAGTAAATAAAATGGCCAAACAGAGAGACAATAAGCGAAATTGCAATTGAGCAATAATT	300
	M G A S P F Y T M N R V I I P F I L P V V L S V I A L N F N S L L T	100
301	CTGACTTCGACTTATCTGATTCCTTTACCATCCCCAGCTCAACCATTAAGTATTACGATTCGATCTGCAGGTGATGAAACAGGCACTCTAATGCA	400
101	GACTGAAGCTCAATAGACATAAGCGAAATGGTAGGGGATCCAGTTGGTAATGCATAATGCTAAGCTAGACGCTCCACTACTTCTGCTGTAGATTAGCTGT	133
	D P D L S V F L Y N P L A Q P L G I T I R S A G D E T A T S N A Q	
401	AGCTCTGGTATTCCTTTATACAAATGCTCTGATGATTATCTGGAACGGTATTATACTTCACACAAAGACCGGGGCTTAAAGTAAGCAAAATAATCATGA	500
134	TGGAACCATTAACCAATATGTTAAACAGACTCTAATAAAGACCTTGCCATAATATGAAGTGCTGTTCTGGCCCCGCACTTCTCATCTCTTTATTAGTACT	164
	A L V F V Y T I V L M I : S G T V L Y F T Q R P G R K V R E .	
501	CAGCEACTAGTCTTGGTTATCAAAATATTGAATAGTTGTGAGGATGTTTATCAGTAGTCACTGGTAGTATAATTTGTTTACAGAGACGGGAGCAAACTC	600
	GTCGGTCACTGAAACCCAATAGTTTATAACTTTATCAACAGTCTTAACAAAATAGTCACTAGTAACCATCATATTAAACCAATCTCTCTCCCTCTTTAG	
601	CCAGCTCGAGGATCCGAACCTATAGTATTTCTCTAGCTGCACTTTGATTATGATGACGAAATGAATACGTTCTTATAAAATTTGGGACAGGAGAT	700
	GTCGGAGCTCGTAGGCTTCAATATCATAAACAAACAGATCGACGTACAACTAATACTACTGCTTACTTATGATAGAAATATTAAACCTGTCTCTTA	
701	CCTACACCATTAGGAGCTCAAGTTATATCAGTGTGGGTTCTAGCGGCTGCAACCATTTCTTATACAGATAAAAAAGAAAAATACAGGCTTGACAACTG	800
	GGATGTGCTAATCTCCGAGTTCAATATAGTCCACACCCAAAAGATCCGCGACCTTGCTAAGAAATAATGTCTATTTTCTTTTAAATGTCAGACTGTGAC	
801	CAGGAGCAATTCGGCTTCGGCAGGAATGGATTAGCTATTGGAGTAGGTTTTATGAGGAGGCTCTTTAGTAGCCATTTCTGTTGGGGTGTGATATC	900
	GTCTCTCCGTAAACCCGAAGCGCTCTTAACTAATCGATAAAGTCACTCAAAAAATAGTCTCTCGAGAAAAATCATCGGTAAAGACAAAACCCCACTATAG	
901	CATGTTCCAAACCACTAAAAAAATATCTGCAAAAATCGTTCTAAAAATGATTGAATGTATATAGTAGTTTAAATCTCTTAG	978
	GTACAGAGTTGGTGATTTTTTATAGACCTTTAGCAAGATTTACTAACTTAAACATATATCATCAATTTAGGAAATC	

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(SEQ ID NO: 47) 1 CAATGTGTTCCCAACTTTTACAAAACATTTCTGAAAAAGAGTTGAAACACTCAAGACCAATTTGTTCAAAATAGGATGGTTGTGTTGATGATG 120
(SEQ ID NO: 48) 1 GTTACACAGGGCTTCAAAAATTTTTGTAGAAGGACTTTTTCTCAAGCTTGTGAGTTTCTGTTAAAGCACTTTATCTTACCAACCACTGATCTAC 120
(SEQ ID NO: 46) 1 M N 2

101 CACAGGATTAGACAAGAGTTGAAAAAGGTGGAGCTGTCTCTACCTACAGAGAGCTTTATGTTCTTTTCCAGGCTTATAGTCAAAAAGCAGTTG 200
CTGTCTAATCTGTTCTCAACTTTTCCCACTTGCACAGCAAGATGGATGTTCTCTGACAAATACCAGAAAAAGGTTCCGGAATCTACTTTTCTGCAAC 200

1 D R I R O E L E K C G A V V L P T T E T V Y G L F S K A L D E K A V D 36

201 ACCATGTTTACCAACTCAAACTGCTCTTACAGATAAGGCACTCAATCTCAATATGCTCTTTCCAGGACCTCTTGCAGCTTTCAAGAATCAGCCAGC 300
TGGTACAAATGGTTGAGTTTGCAGCAGGATCTCTATTCTGCTAGTTAGGTTATAGCGGAGAAAGCTCTGTAGAACGTTGAAAAGTTTCTTAGTGGTCTG 300

3 H V Y Q L K R R P R D K A L N L N I A S F E D I L H F S K N O P A 69

301 TTATCTACAAAACCTTTAGAGAGCTTTTCCAGGTCCTTGCACCTATTTCTGGAAGCCAATGACCGAGTTCCCTATTTGGGTAAATTTCTGACCTTCA 400
AATAGATGTTTCTGACATCTCTGAAAAAGGTTCCAGGGAAGTGGTAATAGAGCTTCCGTTACTGCTCAAGGGAATACCACTTTAAGACTGGAGCT 400

70 Y L O R L V E T F L P G P L T I I L E A N D R V P Y M V N S D L A 102

401 ACTATGGATTTCCGATGCCACTCACCTATCACACTGGATTTAATTCAGAGACAGGTCCTTGATGGGCGCTTCCCAATATCTCAGGTCAAGCA 500
TGATAACCTAAAGCTACGGGTCACTGGGATAGTGTCACTAAATTAAGCTCTCTGTCAGGGAAGTAACCGGCAAGCGTTATAGAGTCCAGTCCGTT 500

103 T I C P R M P S N P : T L D L I R E T G P L I C P S A N I S G Q A S 136

501 GTGGTGAACCTTTGAACAAATTTCAAGGATTTTCAACCAAGAGTTCTGGGTCGGAAGAGCTGCTTTCTAACTGACAGGATTTCACTATTGTGCA 600
CACCACATTTGGAACCTTTTAAAGACTTCTAAAGCTGCTCTCAAGAGCCAGAGCTTTGCTACGAAAAGATTGACCTTGTCTAAGTTGATTAACACT 600

137 G V T F E O I L K D F D O E V L G L E D D A F L T G Q D S T I V D 169

601 TTTGCTGAGACAAGGTTGAAAATTTTACCCAGCGGCAATTAACGAGAAGATATTCTGCTGCTTGCAGAGATTTCTTTTGCAGGAGCTTGAATG 700
AAACAGACTCTGTTTCCACTTTTAGAATGGGTTCCGCTTAATTTGCTCTTATAGAAGAGGCAAGGCTCTCTAAGAAAAGCTCTCCGAACTTAC 700

170 L S G D R V X : L P K A Q L N E K I F L L G C O R F L L R R L E M 202

701 CTAGAGATTTGCAAGAACAGATGTGAAAGCGATATGTGACATCAACCAAGAGCTTTGGGTTATCTTTTACTCCAGAGGAAACCGGTAGCCCACTAG 800
GATTTCTTAAAGCTTTCTTCTACACTTTCCCTATACACTGTATTTGCTTCTCCGAAACCAATATGAAAATCAGGTTCTCTTTGCGGATCGGTTGATC 800

203 L R D L O E T D V K A : C D I N O E A L G Y T F S F E E T A S Q L A 236

801 CTAGACTGTTCTCAGGATTTCCATCAATTTCTTACTTGGCTATGAGATGCAAGTAATCATGTTCTTACTTGGATATGTTCCAGCTGAAGTTTACGAATCACT 900
GATCTGACAGAGCTTTAAGGCTAGTAAAGGATGAACCGATATCTTACTCTGATAGTACAGATGAACCTATACAGGTGGCACTTCAATGCTTATGTA 900

237 R L S O D S M N F L L G Y E D A A N N V L L G Y V M A E V Y E S L 269

901 CTATTCAAAAGCAGGATTTAATATCTTACTTTAGCAGTTTCACTTCAAGGCAAGGTTCAAGGTATCGGTAAAAGTTTACTACAAGGTTGCAACAGAA 1000
GATAAGGTTTCTTCTAAATTTATAGAAATCGTAAGTGGAGTTCCGTTTCCAGTTCCATAGCCATTTTCAATGATGTTCCCAACCTTGTCTTT 1000

270 Y S R A G P N I L A L A V S P O A Q G G I G K S L L O G L E O E 102

1001 GCCAAAAGATGTTGTTATGGGTTTATCCCTTTAAATTTCTGCAATCATGCTCTGGGTCCTCATGCTTTTATGAAAAGTTTGGCTATCTTGTGATAAAA 1100
CGGTTTCTACACCAATACCCAAATAGGCGAATTTAAGACGGTTAGTAGCAGACCCAGCACTACGTAATAATATCTTTTCAACCGATATGAACACTATTTT 1100

303 A K R C G Y C F I R L N S A M N R L G A N A F Y E K V G Y T C D K M 336

1101 TGCAGAAACGGTTTATTCGCACTTTTACTTTGATTTTCTTATTTGTAATAATCAAACTAATGGACTAGTCAACAAATAAGGAGAGACCTATGATTTT 1200
ACGCTTTTCCAAATAGGCTAGAAAATCAAACTAAAGAAATAACATTTTACTTTGATTTACCTGATCAGTGTGTTATTTCTCTTCTGATACTAAAAAC 1200

337 Q K R F I R I P . 345

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[illegible]

- 20 -

1301 TTACTATGCAATTCACCAACAACAAAAAGCTTTCAAAATTAGTCTGTATGACGATCCGACATGGGTACTTTCTGTTGAAAACTCATGAACTCTTTAAAT 1400
AATGATACCTAAACTGCTTTCTTTTTCGGAACTTAAATCAGCACTACTGCTACGCTGTACCCATGAAAGACCACTTTTTCAGTACTTTCAGAAATTA
1354 Y Y G F D E E E K A F E I S R D D D A T W V L S G E R L N K L F N 1386

1401 ATGACCAACTTTGATCGTGATCAATCTGTGATGAACTTTA 1441
TACTGTTTGAAACTAGCACTACTTAGACAGTACTTTGAAAT
1387 M T N F D R D E S V N K L 1399

gsep111

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(SEQ ID NO: 53) : TCGAATGCTTTAAGAAAACATTCAGAAATCAAGAAAAACAGTAAGACAGT.....CTTATGAATTATTAGAAATGAAGAAAGAAAGCATATTAT 100
(SEQ ID NO: 54) : ACCTTACGGGAATTCCTTTGTTAACTTTAGTTCCTTTTCATTCCTCTCAAGAAAAACAGAACTACTTAATAATCTTACTCTCTTTCTTATAATA 1
(SEQ ID NO: 52) : N 1

101 CGCTGAAGAAACAGTAGAACCAAAACCAATTCAGCTTGGTGAATATAAATTTGGTTTCCATGACGATGTAGAGCTGTCTTATCGACAGCAAAAGGACTC 200
CGAAGTCTTTCTCATCTGGTTTGGTTAACTGGAAACCACTTATATTAAACCAAGGTAAGTCTACATCTGGAGCAAGATAGCTGTCTCTTTCTCGAG

2 A E E R V E P K P I D L G E Y K F G F H D D V E P V L S T O R G L 34

201 AAAGAAAGTCTTATTCGTGAATTTATCTGCTCTAAGGCTGAGCTGAGTGGATGTTGGAGTTCCTTTGAAGTCTTATGAAGCTTCAAAAAAATGCCCA 300
TTGCTTCCACAATAAGCACTTAATAGACGAGATTTCCCACTCGGACTCACCTACAACTCAAGGCAAACTTCAGAACTACTTTGGAAGTTTCTTACGGGT

35 N E G V I R E L S A A K G E P E N H L E F R L E S Y E T P R E M P H 40

301 TCGAACTTGGGAGCAGACTTGTGAGAGTTGACTTTGATGACTTAATCTACTACCAAAACCACTGACAAACCAAGCCCTTCTGGGATGATGATCC 400
ACCTTTCAAGCCCTCTCTGAACTCTCTAACTGAACTACTGAAATAGATGATGCTTTTGGTAGACTGTTGGTGGGCAAGAACCTTACTACATGG

69 Q T W G A D L S E I D F D D L I Y Y Q E P S D K P A R S W D D V P 101

401 TGAAGATTAAGAAACCTTTCAAGCTATGGGATTCAGAACTCAAGCTGCTTATTTAGAGGCTCTTCCGCACTACAGTCAAGAGTGGTTTAC 500
ACTTTCTAATTTCTTGAAGCTTGCATAGCCCTAAGCTCTTCACTTGCAGAAATAAATCTGCCCAAGAGCGGTGATCTGCTGCTTCAACAAATG

102 E K I X E T F E R I G I P E A E R A Y L A G A S A Q Y E S E V V Y 134

501 CACAACATGAAGAAAGCTTCCAAAATAGGTATTATCTTTACAGATACAGATTCGGCACTCAAGAAATACCCAGACTTATTTAAACATACTTTGCCA 600
GTCTGTACTCTCTTCAAGGTTTAACTCAATAAGAAATGCTTATGCTCAAGGCTGAGTTCCTTATGGGCTGAAATTAATTTGATGAAGCT

135 N H M K E E F O R L G I I F T D T D S A L K E Y P D L F K Q Y F A K 168

601 ACTTGTACGCGCAGATACCAAGTTGGCAGCTCTCAACTCAGCAGTATGGTGGGCTGAACTTTATCTACGTCGCAAAAGGTGTCAGGTAGATAT 700
TCAACATGGCGCTGTCTATTGTTCAAGCTCGGAGTTGAGTGTCTACACAGCCCACTTGAAGATAGATGACGGTTTCCACAGTTCATCTATA

169 L V P P T D N K L A A L N S A V M S G C T F I Y V P K G V K V D I 201

701 TCACTTCAACTATTTCCTATCAATAACGAAATATAGGTCAGTTCGAAGCTACCTTGATTATCCTTGATGAGGAGCAAGCTCCACTACGTAGAA 800
AGGTGAAGTTGAATAAGGCATAGTTATGCTTTATATCCAGTCAAGCTTGCATGCACTAATAGCAACTACTCCCTGTTGCGAAGTGATGATCTT

202 P L O T Y F R I M N E N : G O F E R T L I I V D E G A S V N Y V E 234

801 CGATGTACAGCACCACATATTCAAGCAATAGCTTACAGCTGCCATTGTAGAAATTTTGGTTTGGAGCGAGCTTATATGCTTATACAACTATCCAAA 900
CTTACATGTCCTGTTGATAGGTTCTTATCGAATGTGGCAGGTAACATCTTTAAAAACGAAACCTGCTTGAATATACGCAATATTTGATAGGTTT

235 G C T A P T Y S S N S L N A A I V E I F A L D G A Y M R Y T T I O N 268

901 ACTGCTGTGATAACCTCTATAACTTGGTAACAAAGCTGCTAAGGCTCAAAAGGATGCCACTGTTGAGTGGATTGATGGAAGCTTGGGTGCCAAACGAC 1000
TGACACAGCTATTGAGATATTGAACCACTGTTTCCGACGATTCGAGTTTCTACGGTGACAACCTCACCTAACTACCTTTGAACCCACGCTTTGCTG

269 M S D N V Y N L V T X R A K A O K D A T V E W I D D N L G A R T T 301

1001 TATGAATATCCATCTTTTACCTTGATGGAGAGGAGCCCTGCTACCACTCTCTATCCCTTTGCTAATGACGGGCAACACCAAGACACGGGTGCT 1100
ATAGTTTATAGGTAGACAAATGGAACTACTCTTCTCGGCAACCATGTTACGAGAGATAGCGGAAACUATTACCTCCCTTGTGTTCTGTCGCCAGCA

302 M K Y P S V Y L D S E G A R C T M L S I A F A M A G O H O D T G A 334

1101 AAGATGATTCACAACTGCTCCACATACAGCTCTGCTATTGTTGCTAAATCCATGCTAAAGGTGAGGAAAGGTTGACTACCTGACAACTCACCTTTA 1200
TTCTACTAAGTGTACGAGGTGATGTTGAGCAGATAACACAGATTTAGGTAGGATTTCCACTCTCTTCCAACTGATGCCACTGTTCAGTGGAAT

335 K M I N M A P N T S S S I V S E S I A R G O C R V D Y R G O V T P N 368

1201 ACAAGAACTTAAGAAATCTTTTCCCACTTGAATGTGATACCATTTATCATGATGACCTT 1263
TGTTCCTGAGATCTTTAGACAAAGGCTGAACCTTACACTATGGAATAGTACCTACTGGA

369 K H S K E S V S N I E C D T I I N D D L 389

99p3362

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(SEQ ID NO: 56) 1 AGCTGCAATTTATGAGCAAGTATCTTATCTTAAAGAAAGGAAGTUTTTATCTAACTCUTTATAATGAAGTTCAAACTGAAACAGCACTTTAATCTTA 100
(SEQ ID NO: 57) 1 TCGACTTAAATACTCTTTCATAGGATAGAAATTTCTTCTTCTTCAAAATAGATTGAGCAATATTACTTCAAGTTTGACTTTTCTGTTGAAATTAGAA 100
(SEQ ID NO: 55) 1 A G I Y E Q V S Y L K E G R S V Y L T R Y N E V Q T E T A T L I L 33

101 GGAGCTATTGTGGGATAGCTAGTTCCTTGTACTCTTTTATCTCTCAATCTTCTATATTTCAGCAATTCGCGGAGATATCTTGATTAAACCAATTT 200
CTCGATTAACACCCCTTATCGATCAAGGAAACAATGAGAAATAGACAGTTAGAGATATAAAGCTCTTAAGGCGGCTCTATAGAACTAATTTGCTTAA 200
34 G A I V C I A S S L L L F Y S V M L L Y P E O P R R D I L I R I S 67

201 CAGCTTTACGATTTTTGAAACACATGCTCAGTATATGTTAGTCAATTTGCCAGTTTGTATTTGCTGCTAGTCTCTTTATTTAAGCAGTCCGAGCTT 300
GTCCAAATGCTAAAAAATCTTGTGTAAGAGTCAATATACCAATCAATTAAGCGGTCAAAACATAAACCAGGATCAGAGAAATAAAATTCCTCAGCTCTGAA 300
68 G L R F F E T H A Q Y M V S O F A S F V F G A S L F I L S R D L 100

301 CGTGATTCGCTTCTCAGCTTTATTAAGTCTTTCTAGCTAGTCCAGTTTTCAGGCTTTACGCTCAAGCGCAGAAAGAAATCTCTGTTTCTATGCAATTTATG 400
CCACTAAGCGAAGGAGTGAATTAATCAGAAAGATCGATCAGCTCAAACTCGGAAATGCGAGTTCCGCTCTTCTTAGAGCAGAAAGATCTGTTAATAC 400
101 V I G L L T L L V F L A S A V L T L Y R Q A O K E S R V S M Y I M 133

401 AAAGCAAAATAGGATGATGCAACTAAAGAATATATCTAAAAAATTTGAGGCGCTCAGCTATTTTCAGATACGAATCTTTA 481
TTTCTTTTCTACTACTACTTGAATTTCTTATATAGATTTTTAAACCTTCGGCAGTCGATAAAAGTCTATGCTTAGAAAT 481
134 K G K 137

9993387

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(SEQ ID NO: 59) : TTTTATCTAGTACAGTATATTTATTCGGCTGTCCCAATATTCAATCCATCCAAATGTTATTAGAAATGGATCTTAGTTTCTTCAAGATATGACGACTGG 100
(SEQ ID NO: 60) : AAAATAGATCATGTATATATAAATAACCGACAGCGTTATTAAGTTAGGTAGGTTTACATAATCTTACCTAGAATCAAAATGAAGTTCTATATCTCTGACC 100
(SEQ ID NO: 58) : M T T G 4

101 AGTATATTGCTTTCCGTTCAATATATATTCCTTTTATTTGATGAATAACTATTTAATAGGTTGGAGTGTGCAATTCGTCTGAATCAATTAAG 200
TCATATAACGAAAGGCAAGTGTATATATAACACAAAAAATAAATACTTATTCATATAAATATCCAACTCACCGGTAAGCAGACTTTAGTTAATTC 200
5 V Y C P P P T Y I L P P P Y L N N Y F N R L S C R I R L K S I K 37

201 CACTTACCAGTTTCTAGTTTCAAATTACGAGCTCTTAGTACGGGATTTGGACGGGCACTTATTTTATTCATTTTCTAATTCGATTAGTAATGGTT 300
GTGAAATGCTCAAAATCAAAATTTAATGCTGAGAATCATGCCCTTAACTCCGCTGAAATAAAAATAACTAAAAAGATTAACTAAATCATTACCAA 300
38 H F T S F S F R L A A L S T G I W T A T L F L L I F L I A F S N G V 71

301 TTAGCTTCTCTTGGAGATAAAGGAGTTCAATTTTAAAGAGATTTTATGATATAAGTATTGCAACAAATGCTAGTTTCTTATAGCAATTTTCTTC 400
AATCGAAGAGAACTCTATTTCTTCAACTAAAGAAATCTCTTAAATACCATATTATACACGTTTGTACCATCAAGAAATATCTTAAAAAAGAG 400
72 S F S L S I K E V D F L R E P Y G I S I A N N A S P F I G P P P S 104

401 TTATATAGCATACTATTTCTTTTATCCTTACTACTATTAGCACTTTTCTTCTGCTTTAAAAAATCAAAATGAGCTTAGTATTTCTGTTTACTTTT 500
AATATATGATATGATAAAGAAAAATAGCAATGAATGATAATGCTCAAAAGAACCAAAATTTTACTTTTGTACTGCAATCATAAAGCAAAATGAAAAAT 500
105 V I A Y Y F P L S L L T I S S F S W P K K S N M S L V F L Y T F L 137

501 TTTGTAGAAATCCTTATCTGGATTTATCAGTTGACAAATGGGATAATGGATTTATTCGCAATTTTCTAGTATATGTTAAATTCCAATCGGTATGCTTCA 600
AAACATCTTAGGAATAGACCTAAATAGTCAACCTGTTACCTTATTAACCTAATAACGGTTAAAAAGTCATATACCAATTAAGGTTAGGCATAGTAACT 600
138 F V E S L F W I Y O L D N G I I G L L P I F O Y N V N S N P Y A L I 171

601 TTTATTCGCTTACATTACTATCTATCATATTCATTGACTGTATTTCTGTTTATAGAACTGGAGGAGAGTGTAAAGTTGGAAATGGGAAAGTTAAG 700
AAATAACGGAATGTATGATAGATAGTATTAGGTAACTGACATAAAGACAACTATCTTTGACCTCTCTCACATTTTCAACCTTTACCTTTCAATTC 700
172 Y W L T L L S I : : P L T V F S V N R N W R R V . 196

[illegible]

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1301 AAGCAATAGCAATCACAAGGTTCTTTTCAAGAAAAATACTATCCAGCTGTAAAGCAAAAGGTTTATCGAACTCGTTTGGCCAAAGGATTGACAGTTTCTT 1400
TTCTTTATCTTACTGTTTCCCAAGAAAACTTTCTTTATGATAGGTCGACATTTCTTTTCCAAATAGCTTGAGCAAAAGGTTTCTTAACTGTCACCA 417
414 C I E *

gop41

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(SEQ ID NO: 65) 1 CTTTTTCACATTTCAAAAGTCTTACGACAGAAAAGAGTCTCTATCTTCAAAAGAAATTTATTTCTTTTCACATCTGCTTTGCTATTTT 100
(SEQ ID NO: 66) CAAAAAAGTGTAAAGTTTTCAGCAATCTGTCTTTCTTTCAGCAGATATCAAGCTTTCTTTTAAATATGGAAGTGTAGACTGAAACCATAAATAA 100

101 TAGAGAAAAATTAAGTTCTCCATGCTTTATGAGAGGTTCTCTTTATGCGAATGAAGATTAGTAGTGGAACTGCGAAATTCAGTCCCAAAAGCACT 200
ATCTCTTTTAAATTCAGAGGCTACCAAAATACCTCTCCAAAGCAAAATACCTTACTTCTAAATCATACCTTAGACCTTTAACTGAGGCTTTTGTCTA 200

(SEQ ID NO: 64) 1 R V Y O E V P V Y A N E D L V V E S O K L T P K T S 26

201 TTTCAATTAACCGAGTCCGCTTAAATTAACAGGAATTCAGTATTTAAGCTATCAATTCATATTTATAGTCCGACAGAAAGATTTTATATGATC 300
AAAGTTTATTTGGCTCAGCGCAATTTATTTGTTCTTAAAGTCTATAATTCATAGTTTATAGTAAATATGAGGCTTTTCTCAAAAATATAGTAG 300

27 F O I T E N R L N K O G I P V P K L S N O F I A A O E R F L Y D Q 60

301 AATCAGAGTAACTCCAGCAATAAAAAGTATGCTTAGAATCTGACTTTAACTGTACAAATAGTCTTATGATTTTAAAGAGAGTGAATTCATCTTATC 400
TTAGTCTCCATTCAGGTTGTTATTTTTCATACCAATCTTAGACTGAAATTTAGCATGTTATCAGCAATAGTAAATTTTCTCACTTAGTAGGAATAG 400

61 S E V T P T I R K V M L E S D P E L Y N S P Y D L R E V K S S L S 93

401 AGCTTATTCGCAAGTATCAATTCAGCAAGCACTGTTTGTAGAGGAAGAGAAATTTCTACATATTCATCAGGCTCGATGCGTAAAGAAATCAACTTCT 500
TCGAATAAGCGTTCATAGTTAGCTGTTCTGTTACAAACATCTCTCTCTCTTAAAGATGTATACTAGTCCGACCTGAGCTTCTTCTTCTTCTTCTTCT 500

94 A Y S O V S I D K T M F V E G R E F L N I D O A O W V A K E S T S 126

501 GAAGAAGATAATCGATGAGTAAAGTTCAGCAATTTATCTCAAAAATATCAGAAAGATTTCTTCTCTATTTATGTTAAGCAACTGACTTCTGGAAG 600
CTTCTCTTATAGCTACTCATTTCAAGTTCTTACAAATAGACTTTTATAGTCTTTCTAAGAAGAGATAAATACAAATTCCTTCACTTAGTAGCTTTCT 600

127 E E D N R N S K V O E N L S E X Y O R D S F S I Y V K O L T T G K E 160

601 AAGTCTGATCAATCAAGATCAAAAGATGTATGCGAGCCAGCTTTTGAAGCTCTCTTATCTCTATTTATACGCAAGAAAAATTAATGAGGCTTTTATCA 700
TTGACCATAGTTAGTTACTTCTTCTTACATACCTGCTCCGCAAACTTTTACAGAAATAGAGATAATATGCTTCTTTTATTTATCTCCAGAAATAGT 700

161 A G I N O D E R N Y A A S V L K L S Y L Y T O E K I N E C L Y O 193

701 GTTAGATACGACTGTAATAATCTCTCTGACTCAATGATTTTCCAGCTCTTATAAACAGAGGCAAGTGTATGCTTCTTCTTAAAAAGAGATAATAAA 800
CAATCTATGCTGACATTTATGCTATAGAGCTCAGTTACTAAAGGCTCAAGAAATTTGCTCTCTTCTCACCATCAGAGAGATTTTCTTCTTATTTT 800

194 L D T T V K Y V S A V N D F P G S Y K P E G S G S L P K K E D N K 226

801 GAATATTTTAAAGGATTTAATACGAAAGTATCAAAAGATCTGATAATGTAGCTCATAATCTATGCGATATTACATTTTCAAAACCAATCTGATGCCA 900
CTTATAAGAAATTTCTTAAATTAATGCTTTCTATAGTTTCTTACACTATTACATGCACTATTAGATAACCTTATAATGTAAGGTTTGTAGACTAGCT 900

227 E Y S L K D L I T K V S K E S D N V A N N L L G Y Y I S N O S D A T 260

901 CATTCAAATCCAGATGCTCTGCAATATGCGAGATCATTCGATCCAAAGAAAAATTCATTTCTTCTAGATGCGCGGCAAGTTTATGGAAGCTATTTA 1000
GTAAGTTTAGGTTCTACAGCGGTAATACCTTCTACTAACCTAGGTTTCTTTTAACTAAAGAGATTCTACCGGCTTCAAAATACCTTGGATAAAT 1000

261 F R S K N S A I N G D D N D P K E K L I S S R N A G R P N E A I Y 293

1001 TAATCAAAATGCAATTTGCTGCTAGAGTCTTCACTAAAACAGATTTTCAATAGTCAGCGAATTCGCAAGGCTGTTCTGTTTAAAGTAGCTCATAAATTCGA 1100
ATTAGTTTACCTAAACAGGATCTCAGAACTGATTTTCTTAAACTATCAGTGGCTTAAAGGTTTCCCAAGAGCAATTTTCACTGATTTTAACTT 1100

294 N C K O F V L E S L T K T D P D S O R I A K C V S V K V A K E I G 326

1101 CATGCGATGAATTAAGCATGATACGCTGTTCTCTATGCGAGATTTCTCATTTATCTTCTTATTTTCACTAAGAAATTCGATTTATGATAGGATTTCTA 1200
CTACGCTCTTAAATTCGATCTATGCGCAACAGATACCTTAAAGAGGTAATAAGAAAGATAAAGTGATTTCTTAAAGCTAATAGTATGCTTAAAGAT 1200

327 C A D E F K N D T G V V Y A D S P F I L S I P T K N S D Y D T I S K 360

1201 ACATAGCCAGGATGTTTATGAGGTTCTAAATGAGCGAACAGATTTTAAATCATTTTCTCAAGAAAGGATATTTTCAAAAGCATCTTAAAGCGGCTT 1300
CTATGCTTCTTCAAAATAGTCCAGATTTTATCTCTTCTTAAAGATTTAGTAAAGAGGTTCTTCTCTTAAAGGTTTCTGATAGGATTTCCGCA 1300

361 : A E S V Y E V L K * 371

106

22

100

63

406

96

388

128

686

263

705

19

80

21

- 28 -

YVES_BACU

(SEQ ID NO: 71) 1 ATGTTAATGCTTTATTCATTATTTGGCTACTTCATAGGCGCATTCATCTGCTTAATTGTGGCGAAGCTTCCAAAGGAATTGATATTGGGAGC 100
(SEQ ID NO: 72) 2 TACAATTAAACCAATAACTAATAAACCACATGAATATCGCTGTAAAGTAGACCGAATTAACACCCGCTTGGAGCGTTTCTTAACATAAGCCCTCG 100
(SEQ ID NO: 70) 1 M L I A L L I I L A Y L I S I P S G L I V O K L A G E D I R E E 34
101 ACCGAAGCGCAACTTAAAGCCTACCAATGCCATTCGTACATTGGGTGTAAGCTGCTTCGCTGTCATAGCCGAGATATTTGAAGGCGACATGGC 200
TCCCTTCGCGCTTGAATCCCGGATGTTTACGTAAAGCATGTAACCCACATTTCCAGCAAGCGCAGTATCGGCTCTATAAACTTTCCTGTGACCG 200
35 G S G M L G A T M A F E T L O V K A G S V V I A G D I L E K T L A 67
201 AACTGCATTGCTTTTCTCATGCAATGATATTCAACCGCTCTTCGAGAGTCTTTGCGGTTTAAAGGCGAGTGTTCCTCATCTTCGCGAATTAA 300
TTGACGTAAACGAAAGAGTACGTACACTATAAGTGGCGAAGACGTCCTCAGAAACGCCAAATCGGTCGCAAAAGGTAGAACGCTTAAATTT 300
68 T A L P F L M H V D I R P L L A G V P A V L G E V P P I F A K F K 100
301 GCGGTAAAGCGGTGGCAGATCAGGAGCGCTTTGCTATTTACGCAACCTGTTATTTATCAGGATGCTTCGCTATTTCTCATCTTTTATCTTGA 400
CGGCAATTTGGCAGCGCTGTAGTCTTCGCAAAAGCATAAATGCGTGGGACAATAAATAGTCTACCAACGCCATAGAGGTAGAAAAATATGAACT 400
101 G G K A V A T S G G V L L F Y A P L L F I T M V A V P P I F L Y L T 134
401 CTAAATTGTTTCTCTCTCATGATGTTAACAGGATCTATCTGTTATATATAGTTTCTTTGTCCATGATAGGTATTTATGATGTTTACCGCTCT 500
GATTTAAACAAAGAGAGATAGCTACAAATGTCCTAGATATGACATATATATCAAGAAACAGGTACTATGCTAAATAACTAACAGCAATGGGACGA 500
135 K P V S L S S M L T G I Y T V I Y S P F V N D T Y L L I V V T L L 167
501 CACTATTTTGTGATATACAGACCGGAGCAATTAAAGCAATTATCAATAAAACAGAACCTAAAGTAAATGTTTATA 582
GTGATAAAAACACTATATGTCGTGCT 582
168 T I F V I Y R N R A N I E R I I N K T E P K V K W L . 193